

GenCore version 5.1.3  
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OM protein - nucleic search, using frame\_plus\_p2n model  
Run on: October 9, 2002, 17:55:09 ; Search time 230 Seconds  
(without alignments)  
6009.201 Million cell updates/sec

Title: US-09-635-501-2  
Perfect score: 4291  
Sequence: 1 MSSSSWLLLSLVAVTAAGT.....ISKGNPNFGQNTDDVQTSF 805

Scoring table: BLCSUM62  
Xgapop 10.0 , Xgapext 0.5  
Ygapop 10.0 , Ygapext 0.5  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 1756436 seqs, 858457221 residues  
Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Command line parameters:  
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-DB=N\_Geneseq\_032802 -QFWT=fastp -SUFFIX=ring -MINMATCH=0.1 -LOOPCL=0  
-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=blsum62 -TRANS=human40.cdi  
-LIST=45 -DOALIGN=200 -THR\_SCORE=pct -THR\_MAX=100 -THR\_MIN=0 -ALIGN=15  
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
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-NO\_XLPXY -NO\_MMAP -LARGEQUERY -NEG\_SCORES=0 -WAIT -LONGLOG -DEV\_TIMEOUT=120  
-WARN\_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7  
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

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pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	4291	100.0	2418	21	AAZ59465	Human MPROT15 codi
2	4291	100.0	3334	22	AAC84366	Human Zace2 protei
3	4291	100.0	3396	21	AAAL2764	CDNA encoding a hu
4	4291	100.0	3396	22	AA02758	Human angiotensin
5	4142	96.5	3732	22	AAS21279	Human CDNA sequenc
6	4061	94.6	2920	22	AAS14880	Human CDNA encodin
7	4013	93.5	2911	22	AAS14890	Human CDNA encodin
8	3740.5	87.2	2262	21	AAZ59466	Human MPROT15 codi
9	3579	83.4	2638	22	AAC84368	Mouse Zace2-5 prot
10	3561	83.0	2638	22	AAC84370	Mouse Zace2-10 pro
11	3509	81.8	2415	22	AAC84367	Human Zace2 protei
12	3119	72.7	3474	22	AAS43515	Human CDNA encodin
13	2899	67.6	2415	22	AAC84369	Mouse Zace2-5 prot
14	1344	31.3	2477	12	AAQ10328	Encodes human test
15	1337	31.2	4020	21	AAA38330	Human angiotensin-
16	1337	31.2	4024	11	AAQ04027	Human angiotensin
17	1337	31.2	4024	20	AAZ35850	Human angiotensin
18	1336	31.1	4024	19	AAV41320	Human angiotensin
19	1334	31.1	3939	22	AAS06085	Angiotensin conver
20	1334	31.1	4563	22	AAS06057	Angiotensin conver
21	1310	30.5	3942	20	AAZ35851	Rat angiotensin co
22	1275	29.7	5005	22	AAH57430	Human intestine ce
23	1086	25.3	2089	23	ABL14379	Drosophila melanog
24	1057	24.6	2074	16	AAQ82948	Tick carboxypeptid
25	1028	24.0	2450	23	ABL16697	Drosophila melanog
26	961	22.4	9006	22	AAH77873	Nucleotide sequenc
27	941.5	21.9	5632	23	ABL14378	Drosophila melanog
28	919.5	21.4	5060	23	ABL16696	Drosophila melanog
29	721	16.8	2082	21	AAZ46692	Degenerate sequenc
30	721	16.8	2082	22	AAZ41469	Human zinc metallo
31	715.5	16.7	1395	22	AAH7876	Nucleotide sequenc
32	502.5	11.7	2025	23	ABL04671	Drosophila melanog
33	478	11.1	313	20	AAV8528	EST clone AU47. H
34	476	11.1	1836	23	ABL27143	Drosophila melanog
35	469	10.9	467	19	AAV09277	Nucleotide sequenc
36	445	10.4	5116	23	ABL04670	Drosophila melanog
37	444	10.3	4001	23	ABL27142	Drosophila melanog
38	414.5	9.7	2046	23	ABL05359	Drosophila melanog
39	387	9.0	666	22	AAF94460	Human hydrophobic
40	387	9.0	1347	22	AAF94470	Human hydrophobic
41	384	8.9	1401	19	AAV40540	Homo sapiens secre
42	383	8.9	847	20	AAZ30083	Human secreted pro
43	381	8.9	848	20	AAZ40770	Secreted protein e
44	381	8.9	848	20	AAZ88191	Human secreted pro
45	381	8.9	848	20	AAZ97564	Extended human sec

ALIGNMENTS

RESULT 1  
AAZ59465  
ID AAZ59465 standard; DNA; 2418 BP.  
XX  
AC AAZ59465;  
XX  
DT 11-APR-2000 (first entry)  
XX  
DE Human MPROT15 coding sequence #1.  
XX  
KW MPROT15; treatment; hypertension; human; myocardial disease; apoplexy;  
KW heart disease; apoplexy; heart disease; nervous denaturation; ds;  
KW Alzheimer's disease; hormone; cytokine.  
XX  
OS Homo sapiens.  
PN JP11318472-A.  
XX

PD 24-NOV-1999.  
 XX PF 22-JAN-1999; 99JP-0014949.  
 XX PR 13-MAY-1998; 98GB-0010373.  
 XX PR 18-AUG-1998; 98GB-0018009.  
 XX (SMIK ) SMITHKLINE BEECHAM PLC.  
 XX WPI: 2000-109268/10.  
 DR P-PSDB; ANY67310.  
 XX MPROT15 polypeptide and MPROT15 polynucleotides - useful for the  
 PT treatment of hypertension, myocardial diseases, apoplexy, heart  
 PT diseases, nervous denaturation, Alzheimer's disease etc.  
 XX Claim 7; Page 14; 22pp; Japanese.  
 CC This is the coding sequence of human MPROT15. The MPROT15 polynucleotide  
 CC and polypeptide sequences can be used for the treatment of hypertension,  
 CC myocardial diseases, apoplexy, heart diseases, nervous denaturation,  
 CC Alzheimer's disease and diseases related to the processing of peptide  
 CC hormones and cytokines.  
 XX SQ Sequence 2418 BP; 744 A; 484 C; 555 G; 635 T; 0 other;  
 Alignment Scores:  
 Pred. No.: 0 Length: 2418  
 Score: 4291.00 Matches: 805  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 100.00% Indels: 0  
 DB: 21 Gaps: 0  
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 QY 1 MetSerSerSerSerTrpLeuLeuSerLeuValAlaValThrAlaAlaGlnSerThr 20  
 DB 1 ATGTCAAGCTCTCCGGCTCCTTCAGCCCTGTGGTGAATCTGCTCAGTCCACG 60  
 QY 21 IleGluGluGlnAlaLysThrPheLeuAspLysPheAsnHisGluAlaGluAspLeuPhe 40  
 DB 61 ATTGAGGACAGCCCAAGACATTTTGGACAGTTTAAACCCAGCCGAGACCTGTC 120  
 QY 41 TyrGlnSerSerLeuAlaSerTrpAsnTyrAsnThrAlaPheLeuGluGlnValGln 60  
 DB 121 TATCAAGTTCACCTTCTGCTTGGATTAACACCAATATTACTGAAGAGATGTC 180  
 QY 61 AsnMetAsnAsnAlaGlyAspLysTrpSerAlaPheLeuLysGluGlnSerThrLeuAla 80  
 DB 181 ACATGCAATTAATGCTGGGACCAATGGTCTGCTCTTTTAAAGGACAGTCCAC 240  
 QY 81 GlnMetTyrProLeuGlnGluLeuGlnAsnLeuThrValLysLeuGlnAlaLeu 100  
 DB 241 CAAATGTATCCACTACAAGAAATTCAGAAATCTCAGATCTCAGCTCAGCTCT 300  
 QY 101 GlnGlnAsnGlnSerSerValLeuSerGluAspLysSerLysArgLeuAsnThrLeu 120  
 DB 301 CAGCAAAATGGGTCTTCAGTGTCTCAGAAAGACAGAGCAACCGTTGAAACAA 360  
 QY 121 AsnThrMetSerThrIleTyrSerThrGlyLysValCysAsnProAspAsnProGlnGlu 140  
 DB 361 AATACANTGAGCACCACATCTACAGTACTGGAAAGTTTGAACCCAGATATCC 420  
 QY 141 CysLeuLeuLeuGluProGlyLeuAsnGluIleMetAlaAsnSerLeuAspTyrAsnGlu 160  
 DB 421 TGTATTATCTTGAACAGGTTTGAATGAATAATGGCAACAGTTAGACTACAA 480  
 QY 161 ArgLeuTrpAlaTrpGluSerTrpArgSerGluValGlyLysGlnLeuArgProLeuTyr 180  
 DB 481 AGGCTCTGGGCTGGGAAAGCTGGAGATCTGAGGTCGCGAAGCAGCTGAGGCC 540  
 QY 181 GluGluTyrValLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAspTyrGly 200

DB 541 GAAGAGTATGTGTCTTGAATAATGAGATGGCAAGAGCAATCATTTATGAGGACTATGG 600  
 QY 201 AspTyrTrpArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGly 220  
 DB 601 GATTATTGGAGAGGAGACTATGAAGTAATGGGTAGATGGCTATGCTACACGCCCGGC 660  
 QY 221 GlnLeuIleGluAspValGluHisThrPheGluGluIleLysProLeuTyrGluHisLeu 240  
 DB 661 CAGTTGATTGAAGATGTCGACATACCTTTGAAGAGATTAAACCATTAATGAACATCT 720  
 QY 241 HisAlaTyrValArgAlaLysLeuMetAsnAlaTyrProSerTyrIleSerProIleGly 260  
 DB 721 CATGCTTATGTGAGGCAAGTTGATGAATGCTATCTCTCTATATCATGTCGAATGGA 780  
 QY 261 CysLeuProAlaHisLeuLeuGlyAspMetTrpGlyArgPheTrpThrAsnLeuTyrSer 280  
 DB 781 TGCTCTCCCTGCTCATTTGCTGTGATATGGGTAGATTGGGACAAATCTGTACTCT 840  
 QY 281 LeuThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGln 300  
 DB 841 TTGACAGTTCCCTTTGGACAGAAACCAACATAGATGTTACTGATGCAATGCTGGACCAG 900  
 QY 301 AlaTrpAspAlaGlnArgIlePheLysGluAlaGluLysPhePheValSerValGlyLeu 320  
 DB 901 GCCTGGGATGTCACAGAGAAATATCAAGGAGGCGGAGAGTTCTTTGATCTGTGTCT 960  
 QY 321 ProAsnMetThrGlnGlyPheTrpGluAsnSerMetLeuThrAspProGlyAsnValGln 340  
 DB 961 CCTAATATGACTCAAGGATCTGGGAAATTCATCTAAGCGACCCAGGAATGTTCCAG 1020  
 QY 341 LysAlaValCysHisProThrAlaTrpAspLeuGlyLysGlyAspPheArgIleLeuMet 360  
 DB 1021 AAAGCAGTCTGCCATCCACAGCTTGGGACCTGGGGAAGGCGACTTCAGGATCTTATG 1080  
 QY 361 CysThrLysValThrMetAspAspPheLeuThrAlaHisGluMetGlyHisIleGln 380  
 DB 1081 TGCACAAAGTGACAAATGAGGAGCTTCTTCAGAGCTCATCATGAGATGGGCATATCCAG 1140  
 QY 381 TyrAspMetAlaTyrAlaAlaGlnPropheLeuLeuArgAsnGlyAlaAsnGluGlyPhe 400  
 DB 1141 TATGATATGGCATATGCTGCACAACTTTCTGCTAAGAAATGGAGCTAATGAAGATTC 1200  
 QY 401 HisGluAlaValGlyLysLeuMetSerLeuSerLeuAlaThrProLysHisLeuLysSer 420  
 DB 1201 CATGAAGCTGTGGGAAATCATGTCTCTGAGCCACACCTTAAGCATTTAAATCC 1260  
 QY 421 IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeuLeu 440  
 DB 1261 ATTGCTCTCTGTCACCCGATTTTCAGAGAGACAAATGAACAGAAATAAATTCCTGCTC 1320  
 QY 441 LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTrpArg 460  
 DB 1321 AAACAGCACTCAGCATTTGGGACTCTGCCATTTACTTACATGTTAGAGAGTGGAGG 1380  
 QY 461 TrpMetValPheLysGlyGluIleProLysAspGlnTrpMetLysLysTrpTrpGluMet 480  
 DB 1381 TGTATGCTTTTAAAGGGAAATTCCTCAAGAGCAGTGGATGAAAGTGGTGGGAGATG 1440  
 QY 481 LysArgGluIleValGlyValGluProValProHisAspGluThrTyrCysAspPro 500  
 DB 1441 AAGCGAGACATAGTTGGGCTGCTGGAACTGTCGCCCATGATGAACATCTGTCGACCCC 1500  
 QY 501 AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrTrpThrArgThrLeu 520  
 DB 1501 GCATCTCTCTCCATGTTTCTAATGATTAATCTCATCTCTGATATTACAAAGGACCCCT 1560  
 QY 521 TyrGlnPheGlnPheGlnGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis 540  
 DB 1561 TACCAATTCAGTTTCAAGAGCAGCTTTGTCAGAGCAGCTTAACATGTAAGGCCCTCTG 1620  
 QY 541 LysCysAspIleSerAsnSerThrGluAlaGlyGlnLysLeuPheAsnMetLeuArgLeu 560

CDS 35..2452  
 FT /\*tag= a  
 FT /product= "Zace2"  
 XX  
 PN WO200070032-A1.  
 XX  
 PD 23-NOV-2000.  
 XX  
 PF 03-MAY-2000; 2000WO-US11932.  
 XX  
 PR 13-MAY-1999; 99US-0311482.  
 PR 27-AUG-1999; 99US-0384706.  
 XX  
 PA (ZYMO ) ZYMOGENETICS INC.  
 XX  
 PI Piddington CS, Petrie CR, Shoemaker KE, Bishop PD;  
 DR WPI; 2001-025018/03.  
 DR P-PSDB; AAB48095.  
 XX  
 PT Angiotensin-converting enzyme, Zace2, useful for treating inflammatory  
 PT bowel disease, e.g. Crohn's disease and ulcerative colitis, or diseases  
 PT associated with inflammation such as arthritis and enterocolitis -  
 XX  
 PS Example 1; Page 95-100; 125pp; English.  
 XX  
 CC The invention relates to the metalloenzyme Zace2. Zace2, an angiotensin-  
 CC converting enzyme is a zinc metalloproteinase that plays roles in blood  
 CC pressure regulation and fertility. Zace2 can be expressed by standard  
 CC recombinant methodology. Zace2 polypeptides are useful for treating an  
 CC inflammatory bowel disease (e.g. Crohn's disease and ulcerative colitis),  
 CC diseases associated with inflammation like arthritis and enterocolitis,  
 CC as targets for identifying modulators of zinc protease activity, for  
 CC screening or identifying new angiotensin-converting enzyme (ACE)  
 CC inhibitors, and as a basis for rational drug design for inhibitory  
 CC molecules. The nucleic acids can be used to detect the expression of a  
 CC zace2 gene in a biological sample, as probes for in vivo diagnosis and  
 CC for detecting and localizing Zace2 gene expression in tissue samples,  
 CC to determine whether a subject's chromosomes contain a mutation in the  
 CC Zace2 gene, and to detect aberrations associated with the Zace2 locus.  
 CC Inhibitors of ACE are used for treating hypertension of various  
 CC conditions, including left ventricular systolic dysfunction, progressive  
 CC renal impairment, scleroderma renal crisis, congestive heart failure due  
 CC to dysfunction, and treatment of atherosclerosis. Zace2 agonists may be  
 CC used to treat infertility while Zace2 antagonists are used for inducing  
 CC infertility. The present sequence represents a cDNA encoding the human  
 CC Zace2 protein.  
 XX  
 SQ Sequence 3334 BP; 1011 A; 640 C; 754 G; 929 T; 0 other;  
 XX  
 Alignment Scores:  
 Pred. No.: 0 Length: 3334  
 Score: 4291.00 Matches: 805  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 100.00% Indels: 0  
 DB: 22 Gaps: 0  
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 Oy 1 MetSerSerSerTrpLeuLeuLeuSerLeuValAlaValThrAlaLaGlnSerThr 20  
 Db 35 ATGTCAAGCTCTTCTGGCTCTCTTCACGCTGTGTCTACTGCTGCTAGCTCCACC 94  
 Oy 21 lLeGluGluGlnAlaLysThrPheLeuAspLysPheAsnHisGluLaGluAspLeuPhe 40  
 Db 95 ATTGAGGAACAGCCCAAGACATTTTGGACAACCTTTAACCCACCAAGCCGAACCTGTC 154  
 Oy 41 TyrGlnSerSerLeuAlaSerTrpAsnTyrAsnThrAsnIleThrGluGluAsnValGln 60  
 Db 155 TATCAAAAGTTCACTGCTCTTCTGGAAATTATACACCAATATTACTGAAGAATGTCCAA 214  
 Oy 61 AsnMetAsnAsnAlaGluValAspLysTrpSerAlaPheLeuLysGluGlnSerThrLeuAla 80

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Db 215 AACATGAATAATGCTGGGGACAAATGGTCTGCTCTTTTAAAGAAACAGTCCACACTTGGC 274  
Qy 81 GlnMetTyrProLeuGlnGluIleGlnAsnLeuThrValLysLeuGlnLeuAlaLeu 100  
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Db 275 CAAATGATCCACTACAGAAATTCAGAAATCTCACAGTCAAGCTTCAGCTGAGGCTCTT 334  
Qy 101 GlnGlnAsnGlySerSerValLeuSerGluAspLysSerLysArgLeuAsnThrIleLeu 120  
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Db 335 CAGCAAAATGGGTCTTCAGTGTCTCAGAAAGACAGAGCAACGGTTGAACACAAATCTTA 394  
Qy 121 AsnThrMetSerThrIleTyrSerThrGlyLysValCysAsnProAspAsnProGlnGlu 140  
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Db 395 AATACAAATGAGCAACCATCTACAGTACTGGAAAAGTTGTAAACCCAGATATCCACAAGAA 454  
Qy 141 CysLeuLeuLeuProGlyLeuAsnGluIleMetAlaAsnSerLeuAspTyrAsnGlu 160  
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Db 455 TGCCTATTACTTGAACCCAGTTTGAATGAATAATGGCAACAGTTTACACTACAAATGAG 514  
Qy 161 ArgLeuTrpAlaTrpGluSerTrpArgSerGluValGlyLysGlnLeuArgProLeuTyr 180  
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Db 515 AGGCTCTGGGCTTGGGAAAGCTGGAGATCTGAGTGGCAAGCAGCTGAGGCCATTATAT 574  
Qy 181 GluGluTyrValLeuLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAspTyrGly 200  
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Db 575 GAAGAGTATGTGCTCTGAAAATAGATGGCAAGAGCAAAATCAATTATGAGGACTATGGG 634  
Qy 201 AspTyrTrpArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGly 220  
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Db 635 GATTATTGGAGAGGAGACTATGAAGTAATGGGTAGATGGCTATGACTACAGCCGCGG 694  
Qy 221 GlnLeuIleGluAspValGluHisThrPheGluGluIleLysProLeuTyrGluHisLeu 240  
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Db 695 CAGTTGATTGAAGATGTGGAAACATACCTTTGAAGAGATTAAACCAATTATATGAACATCTT 754  
Qy 241 HisAlaTyrValArgAlaLysLeuMetAsnAlaTyrProSerTyrIleSerProIleGly 260  
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Db 755 CATGCCATATGGGCAAGTGTGATGAATGATGATGCTATCTCTCTATATCATCATCTTA 814  
Qy 261 CysLeuProAlaHisLeuLeuGlyAspMetTrpGlyArgPheTrpThrAsnLeuTyrSer 280  
|||||  
Db 815 TGGCTCCCTGCTCATTTGCTTGGTGTATGTGGGGTAGATTTTGGCAAAATCTGTACTCT 874  
Qy 281 LeuThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGln 300  
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Db 875 TTGACAGTTCCCTTTGGACAGAAACCAACATAGATGTACTGTATGATGCAATGGTGACAG 934  
Qy 301 AlaTrpAspAlaGlnArgIlePheLysGluAlaGluLysPhePheValSerValGlyLeu 320  
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Db 935 GCCTGGGATGCACAGAGAAATATCAAGGAGGCGGAGAAAGTTCTTTGTATCTGTGGTCTT 994  
Qy 321 ProAsnMetThrGlnGlyPheTrpGluAsnSerMetLeuThrAspProGlyAsnValGln 340  
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Db 995 CCTAATATGACTCAAGGATCTGGGAAAATTCATGCTAACGGAGCCAGAAATGTTTACG 1054  
Qy 341 LysAlaValCysHisProThrAlaTrpAspLeuGlyLysGlyAspPheArgIleLeuMet 360  
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Db 1055 AAAGCAGTCTGCCATCCACAGCTTGGGACCTGGGCAAGGGGAGCTTCAGGATCCATTATG 1114  
Qy 361 CysThrLysValThrMetAspAspPheLeuThrAlaHisHisGluMetGlyHisIleGln 380  
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Db 1115 TGCACAAAGGTGACAAATGACGACTTCTTCACAGCTCATCATGAGATGGGCATATCCAG 1174  
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Db 1175 TATGATATGTCATATGCTGCACAACTTTCTGCTAAGAAATGGAGCTAATGAAGATTC 1234  
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Db 1235 CATGAAGCTGTTGGGAAATCATGTCTACCTTTCTGACGCCACACCTAAGCATTTAAAAATCC 1294  
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Qy 441 LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTrpArg 460  
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Db 1355 AAACAAGCACTCACCATTGTTGGGACTTGGCCATTTACTTACATGTTAGAGAAGTGAGG 1414  
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Db 1415 TGGATGGTCTTTAAAGGGGAATTTCCAAAAGACCAGTGGATGAAAAGTGTGGGAGATG 1474  
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Qy 501 AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrThrArgThrLeu 520  
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Qy 521 TyrGlnPheGlnPheGlnGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis 540  
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Db 1655 AATGTGACATCTCAAACTCTACAGAGCTGGACAGAACTGTTCAATATGCTGAGGCTT 1714  
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Db 1715 GCAAAATCAGAACCCCTGGACCTAGCATTTGGAATAATGTTGTAGGAGCAAGAACATGAAT 1774  
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Qy 721 SerLeuGluPheLeuGlyIleGlnProThrLeuGlyProProAsnGlnProProValSer 740  
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QY 361 CysThrLysValThrMetAspPheLeuThrAlaHisHisGluMetGlyHisIleLeu 380
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QY 401 HisGluAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLysHisLeuLysSer 420
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QY 521 TyrGlnPheGlnPheGlnGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis 540
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DB 2122 TTTAATTTCTTGTCTACTGCACCTAAAATGTGCTGTGATATCTTCTAGAACTGAAGTT 2181
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RESULT 4
AAD02758
ID AAD02758 standard; cDNA; 3396 BP.
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AC AAD02758;
XX
DT 31-MAY-2001 (first entry)
XX
DE Human angiotensin converting enzyme-2 (ACE-2) cDNA.
XX
KW Human; angiotensin converting enzyme-2; ACE-2; peptidyl dipeptidase A;
KW inflammation; pain; ss.
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OS Homo sapiens.
FH
FT Key
FT CDS
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FT mat_peptide
FT /*tag= c
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PN US6194556-B1.
XX
PD 27-FEB-2001.
XX
PF 11-DEC-1997; 97US-0989299.
XX
PR 11-DEC-1997; 97US-0989299.
XX
PA (MILL-) MILLENNIUM PHARM INC.
XX
PI Acton SL, Robison KE;
XX
XX WPI; 2001-210604/21.
DR
P-PSDB; AAY72667.
XX
XX Novel genes encoding angiotensin converting enzyme-2 useful as
XX antiseize or antigenic agents for therapeutics, diagnostics and
XX screening assays -
XX
XX Claim 1; Fig 1; 76pp; English.
XX
XX The present sequence is human angiotensin converting enzyme-2 (ACE-2)
XX
```

CC cDNA. ACE is also referred as peptidyl dipeptidase A. Nucleic acid  
 CC sequence encoding ACE-2 is useful as antisense or in the analysis of  
 CC sequence specific modulation of gene expression or in the analysis of  
 CC single base-pair mutations in the gene. Nucleic acid sequence encoding  
 CC ACE-2 is useful in therapeutics, diagnostics and in screening assays.  
 CC ACE-2 antagonist is used to treat hypertension or congestive heart  
 CC failure (CHF). ACE agonist is used to reduce the inflammation and pain  
 CC resulting from an insect sting or bite, which was accompanied by an  
 CC injection of bradykinin. Anti-ACE-2 antibodies are used to monitor ACE-2  
 CC protein levels for determining the disease or condition associated with  
 CC an aberrant protein level.  
 XX

SQ Sequence 3396 BP; 1034 A; 659 C; 772 G; 931 T; 0 other;

Alignment Scores:

Pred. No.:	0	Length:	3396
Score:	4291.00	Matches:	805
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	100.00%	Indels:	0
DB:	22	Gaps:	0

US-09-635-501-2 (1-805) x AAD02758 (1-3396)

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Qy	21	IleGluGluGlnAlaIleLysThrPheLeuAspLysPheAsnHisGluAlaGluAspLeuPhe	40
Db	142	ATGTAGGACAGGCGCAAGACATTTTGGACAAAGTTTAAACACGAGCGCGAAGACCTGTC	201
Qy	41	TyrGlnSerSerLeuAlaSerTrpAsnTyrAsnThrAsnIleThrGluGluAsnValGln	60
Db	202	TATCAAGTTCACCTGCTCTGGAATTATACACCAATATTACTGAGAGAAATGTCGAA	261
Qy	61	AsnMetAsnAlaGlyAspLysTrpSerAlaPheLeuLysGluGlnSerThrLeuAla	80
Db	262	AACATGAATAATGCTGGGACAAATGGTCTGCTCTTTAAAGAAACAGTCCACACTGGC	321
Qy	81	GlnMetTyrProLeuGlnGluIleGlnAsnLeuThrValLysLeuGlnLeuAlaLeu	100
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Qy	161	ArgLeuTrpAlaTrpGluSerTrpArgSerGluValGlyLysGlnLeuArgProLeuTyr	180
Db	562	AGGCTCTGGGCTTGGGAAAGCTGGAGATCTGAGGTCGGCAAGCAGCTGAGGCCATTAT	621
Qy	181	GluGluTyrValValLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAspTyrGly	200
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Qy	201	AspTyrTrpArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGly	220
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Qy	261	CysLeuProAlaHisLeuLeuGlyAspMetTrpGlyArgPheTrpThrAsnLeuTyrSer	280
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Qy	281	LeuThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGln	300
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Qy	301	AlaTrpAspAlaGlnArgIlePheLysGluAlaGluLysPhePheValSerValGlyLeu	320
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Qy	321	ProAsnMetThrGlnGlyPheTrpGluAsnSerMetLeuThrAspProGlyAsnValGln	340
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Qy	341	LysAlaValCysHisProThrAlaTrpAspLeuGlyLysGlyAspPheArgIleLeuMet	360
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Qy	361	CysThrLysValThrMetAspAspPheLeuThrAlaHisGluMetGlyHisIleGln	380
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Qy	381	TyrAspMetAlaTyrAlaIleGlnProPheLeuLeuArgAsnGlyAlaAsnGluGlyPhe	400
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Qy	401	HisGluAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLysHisLeuLysSer	420
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Qy	421	IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeu	440
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Qy	441	LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGlyLysTrpArg	460
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Qy	461	TrpMetValPheLysGlyGluIleProLysAspGlnTrpMetLysLysTrpTrpGluMet	480
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Qy	481	LysArgGluIleValGlyValValGluProValProHisAspGluThrTyrCysAspPro	500
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Qy	501	AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrThrArgThrLeu	520
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Db	1822	GTAAGGCCACTGCTCAACTACTTTGAGCCCTTATTTACCTGGCTGAAGACCAACAAG	1881
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 DT 24-OCT-2001 (first entry)  
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 DE Human cDNA sequence encoding for PRO1885 polypeptide.  
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 KW Human secretory and transmembrane; PRO; mammalian; cancer; lung;  
 KW breast; prostate; cervical; tumour necrosis factor-alpha; TNF-alpha;  
 KW cartilage; ear; proliferation; glucose; free fatty acid; skeletal muscle;  
 KW adipocyte; A-peptide; factor VIIA; gene therapy; ss.  
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 OS Homo sapiens.  
 XX  
 PN WO20010466-A2.  
 XX  
 PD 07-JUN-2001.  
 XX  
 PF 01-DEC-2000; 2000WO-US32678.  
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 PR 01-DEC-1999; 99WO-US28301.  
 PR 01-DEC-1999; 99WO-US28634.  
 PR 02-DEC-1999; 99WO-US28551.  
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 PR 09-DEC-1999; 99US-0170262.  
 PR 16-DEC-1999; 99WO-US30095.  
 PR 20-DEC-1999; 99WO-US30911.

PR 20-DEC-1999; 99WO-US30999.  
 PR 30-DEC-1999; 99WO-US31243.  
 PR 06-JAN-2000; 2000WO-US00277.  
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 PR 11-FEB-2000; 2000WO-US03565.  
 PR 18-FEB-2000; 2000WO-US04341.  
 PR 18-FEB-2000; 2000WO-US04342.  
 PR 22-FEB-2000; 2000WO-US04414.  
 PR 24-FEB-2000; 2000WO-US04914.  
 PR 24-FEB-2000; 2000WO-US05004.  
 PR 01-MAR-2000; 2000WO-US05601.  
 PR 20-MAR-2000; 2000WO-US07377.  
 PR 21-MAR-2000; 2000WO-US07532.  
 PR 30-MAR-2000; 2000WO-US08439.  
 PR 17-MAY-2000; 2000WO-US13705.  
 PR 22-MAY-2000; 2000WO-US14042.  
 PR 30-MAY-2000; 2000WO-US14941.  
 PR 02-JUN-2000; 2000WO-US15264.  
 PR 10-NOV-2000; 2000WO-US30873.  
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 PA (GETH ) GENENTECH INC.  
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 PI Baker KP, Beresini M, Deforge L, Desnoyers L, Filvaroff E, Gao W;  
 Gerritsen ME, Goddard A, Godowski PJ, Gurney AL, Sherwood S;  
 PI Smith V, Stewart TA, Tumas D, Watanabe CK, Wood WI, Zhang Z;  
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 DR WPI: 2001-408281/43.  
 P-PSDB: AAU12207.  
 XX  
 PT Isolated, secretory and transmembrane PRO polypeptide used to detect  
 other PRO polypeptides, link bioactive molecules to cells expressing  
 PRO polypeptides, and detect the presence of mammalian tumours e.g.  
 PT lung, breast, prostate, cervical -  
 XX  
 PS Claim 3; Fig 71; 813pp; English.  
 XX  
 CC AAS21244-AAS21518 encode for novel human secretory and transmembrane  
 PRO polypeptides. The PRO polypeptides are useful to detect other  
 CC PRO polypeptides, to link bioactive molecules to cells expressing  
 CC PRO polypeptides, to modulate biological activities of cells expressing  
 CC PRO polypeptides, and to detect the presence of mammalian lung, colon,  
 CC breast, prostate, rectal, cervical or liver tumours by comparing PRO  
 CC polypeptide expression in a cell sample to that in a control sample.  
 CC Some of the 275 sequences are also useful to stimulate the release of  
 CC tumour necrosis factor-alpha (TNF-alpha) from human blood, the  
 CC proliferation or differentiation of chondrocytes, the proliferation or  
 CC gene expression in pericyte cells, the release of proteoglycans from  
 CC cartilage, the proliferation of inner ear utricular supporting cells or  
 CC of T-lymphocytes, the release of a cytokine from peripheral blood  
 CC monocytes (PBMCs), or the proliferation of endothelial cells. Some of  
 CC the PRO polypeptides may modulate glucose or free fatty acid uptake by  
 CC skeletal muscle cells or by adipocytes; or inhibit binding of A-peptide  
 CC to factor VIIA. The PRO polypeptides can be used in assays to identify  
 CC molecules involved in binding interactions. The polynucleotides encoding  
 CC PRO polypeptides can be used to generate probes, antisense RNA/DNA,  
 CC transgenic or knock out animals and can be used in gene therapy.  
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 SQ Sequence 3732 BP; 1137 A; 722 C; 821 G; 1052 T; 0 other;  
 Alignment Scores:  
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 Score: 4142.00 Matches: 802  
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 Best Local Similarity: 85.23% Mismatches: 2  
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 (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||) (|||||)  
 Db 40 ATGTCAGGCTCTTCCCTGGGCTCTTCTCAGGCTTGTGCTAACTGCTCTCAGTCCACC 99



QY 21 IleGluGluGlnAlaLysThrPheLeuAspLysPheAsnHisGluAlaGluAspLeuPhe 40  
 DB 100 ATTGAGAACAGCCCAAGACATTTTGGACAAGTTTAAACCACGAGCCGAGACCTGTC 159  
 QY 41 TyrGlnSerSerLeuAlaSerTrpAsnThrAsnThrAsnIleThrGluGluAsnValGln 60  
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 QY 201 AspTyrTrpArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGly 220  
 DB 640 GATATTCGAGAGGAGACTAGAGTAATGGGTAGATGGCTATGACTACAGCCGCGGC 699  
 QY 221 GlnLeuIleGluAspValGluHisThrPheGluGluIleLysProLeuTyrGluHisLeu 240  
 DB 700 CAGTTGATTGAAGATGTGGAACATACCTTTGAAGAGATTAAGACATATATGAACATCT 759  
 QY 241 HisAlaTyrValArgAlaLysLeuMetAsnAlaTyrProSerTyrIleSerProIleGly 260  
 DB 760 CATGCCATATGTGAGGGCAAGATGTATGAATGCCATATCTCTATATATCAGTCCAAATGGA 819  
 QY 261 CysLeuProAlaHisLeuLeuGlyAspMetTrpGlyArgPheTrpThrAsnLeuTyrSer 280  
 DB 820 TGCCTCCCTGCTCATTTGCTGGTGATATGTGGGTAGATTTTGGACAAATCTGTACTCT 879  
 QY 281 LeuThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGln 300  
 DB 880 TTGACAGTTCCTCTTGGACAGAAACCAACATAGATGTTACTGATGCAATGGTGACACC 939  
 QY 301 AlaTrpAspAlaGlnArgIlePheLysGluAlaGluLysPhePheValSerValGlyLeu 320  
 DB 940 GCCCTGGATGCACAGAGAAATATTCAAGAGGCGGAGAGTCTTTGCTATCTGTGGTCTT 999  
 QY 321 ProAsnMetThrGlnGlyPheTrpGluAsnSerMetLeuThrAspProGlyAsnValGln 340  
 DB 1000 CCTAATATGACTCAAGGATTTGGGAAATTTCCATGCTAAACGGACCCAGGAAATGTTCCAG 1059  
 QY 341 LysAlaValCysHisProThrAlaTrpAspLeuGlyLysGlyAspPheArgIleLeuMet 360  
 DB 1060 AAAGCAGTCTGCCATCCACAGCTTGGACCTGGGGAAGGCGGAGTTCAGGATCTCTTATG 1119  
 QY 361 CysThrLysValThrMetAspPheLeuThrAlaHisHisGluMetGlyHisIleGln 380  
 DB 1120 TGCACAAAGGTGCACAAATGGAGCCTCTCTCAGCTCATCATGATGAGATGGGCATATCCAG 1179  
 QY 381 TyrAspMetAlaTyrAlaAlaGlnProPheLeuLeuArgAsnGlyAlaAsnGluGlyPhe 400

DB 1180 TATGATATGGCATATGCTGCACAACTTTTCTGCTAAGAAATGGAGCTAATGAAGGATTC 1239  
 QY 401 HisGluAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLysHisLeuLysSer 420  
 DB 1240 CATGAAGCTGTTGGGGAATCATGTACATTTCTGAGCCACACCTTAAGCATTTAAATCC 1299  
 QY 421 IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeuLeu 440  
 DB 1300 ATTTGCTCTCTGCTACCCGATTTTCAAGAAGACAATGAACAGAAATAAATTCCTGCTC 1359  
 QY 441 LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTrpArg 460  
 DB 1360 AAACAAGCACTCACGATTTGTTGGACTCTGCCATTTACTTACATGTTAGAGAAGTGGAG 1419  
 QY 461 TrpMetValPheLysGlyGluIleProLysAspGlnTyrMetLysLysTrpTrpGluMet 480  
 DB 1420 TGGATGGTCTTTAAAGGGGAAATTTCCCAAGACCACTGGATGAAAAGTGGTGGGAGATG 1479  
 QY 481 LysArgGluIleValGlyValValGluProValProHisAspGluThrTyrCysAspPro 500  
 DB 1480 AAGCGAGAGATAGTTGGGTGGTGGAACTGTGCCCATGATGAACATATCTGTGACCC 1539  
 QY 501 AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrTyrThrArgThrLeu 520  
 DB 1540 GCATCTCTGTTCCATGTTTCTGATGATTACTCATTTCAATTCGATATTACAAAGGACCTT 1599  
 QY 521 TyrGlnPheGlnPheGlnAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis 540  
 DB 1600 TACCAATTTCCAGTTTCAGAGACACTTTGTCAGCAGACTAAACATGAAGGCCCTCTGCAC 1659  
 QY 541 LysCysAspIleSerAsnSerThrGluAlaGlyGlnLysLeuPhe 555  
 DB 1660 AAATGTGACATCTCAAACTCTACAGAAGCTGCAGAGAACTGTT-GTAAGAAATACCTCA 1718  
 QY 555 ----- 555  
 DB 1719 AAATGTTGAACCTTCCTAGTATTGATTCAGTATTACTCATTTTCCATGCTAGGTTGTATTG 1778  
 QY 555 ----- 555  
 DB 1779 ATTTCTTTGTTCTAAAAAGAAAAATTTTATGGCTCAAAATGCTCCTCATTTACAAACCAA 1838  
 QY 555 ----- 555  
 DB 1839 CATTTAAATTTGTGTGTCAGACAGAACTTAGACCATACAAATTTGGTGGGCCACCTTT 1898  
 QY 555 ----- 555  
 DB 1899 TTCTCCCTATCATAACTACAGCCCTCTCTTCTGGTAATTTGGAAGAAAGACGGTTTGTAG 1958  
 QY 555 ----- 555  
 DB 1959 GGTGGAATATATCTCTTAATATGATCTCTTTTCTTATCTGCCAGAAGCAAAATTTAGCCAA 2018  
 QY 555 ----- 555  
 DB 2019 GTCAAAGAGAAACCATAGATCATAGATGTAATATATATGATCTGTGAACCCCTCAA 2078  
 QY 556 ----- AsnMetLeuArgLeuGlyLysSerGluPro 565  
 DB 2079 AAGGCCCTGAACCCCTTTTGTGTAGCAATATGCTGAGGCTTGGAAAAATCAGAACCC 2138  
 QY 566 TrpThrLeuAlaLeuGluAsnValValGlyAlaLysAsnMetAsnValArgProLeuLeu 585  
 DB 2139 TGGACCCCTAGCATTTGAAAAATGTTGTAGCAGC-AAGAACAATGAATGAAGGCCACTGCTC 2197  
 QY 586 AsnTyrPheGluProLeuPheThrTrpLeuLysAspGlnAsnLysAsnSerPheValGly 605  
 DB 2198 AACTACTTGTAGCCCTTATTACCTGGCTGAAAGACCAAGAAATTTCTTTTGTGGGA 2257  
 QY 606 TrpSerThrAspTrpSerProTyrAlaAsp-GlnSerIleLysValArgIleSerLeuLeu 625

Db 2258 TGGAGTACCGACTGGAGTCCATATGTCAGACCCAAAGCATCAAAAGTGAGGATAAGCCATAA 2317  
QY 525 SSeRAlaLeuGlyAspLySaLaTyrrGluTrpAsnAspAsnGluMetTyrrLeuPheArgse 645  
Db 2318 ATCAGCTCTGGAGATAAAGCATATGAATGGAACGACAAATGAATGATCTGTTCGATC 2377  
QY 645 rSerValAlaTyrrAlaMetArgGlnTyrrPheLeuLysValLysAsnGlnMetIleLeuPh 665  
Db 2378 ATCTGTTGCATATGCTATGAGCGAGTACTTTTAAAGTAAAAATCAGATGATCTTTT 2437  
QY 665 eGlyGluGluAspValArgValAlaAsnLeuLysProArgIleSerPheAsnPhePheVa 685  
Db 2438 TGGGGAGGAGGATGTGCGAGTGCTAATTTGAAACCAAGAAATCTCCTTAATTTCTTGT 2497  
QY 685 lThrAlaProLysAsnValSerAspIleIleProArgThrGluValGluLysAlaIleAr 705  
Db 2498 CACTGCACCTAAAATGTGCTGATATCATCTCTAGAACTGAAAGTGAAGGCCATCAG 2557  
QY 705 gMetSerArgSerArgIleAsnAspAlaPheArgLeuAsnAspAsnSerLeuGluPheLe 725  
Db 2558 GATGTCCCGAGCGGTATCAATGATGCTTTCCGCTCTGAATGACACAGCCCTAGAGTTCT 2617  
QY 725 uGlyIleGlnProThrLeuGlyProProAsnGlnProProValSerIleTrpLeuIleVa 745  
Db 2618 GGGGATACACCCACACTTGGACCTCTTAACACAGCCCTGTTTCCATATGCGTGATGT 2677  
QY 745 lPheGlyValValMetGlyValIleValValGlyIleValIleLeuIlePheThrGlyI 765  
Db 2678 TTTTGGAGTGTGATGGGAGTGATAGTGTGGCATGTGCATCTGATCTTCATCGGAT 2737  
QY 765 eArgAspArgLysLysAsnLysAlaArgSerGlyGluAsnProTyrAlaSerIleAs 785  
Db 2738 CAGAGATCGGAAGAGAAAAATAAGACAGAGTGGAGAAAAATCCTATGCCCTCCATCGA 2797  
QY 785 pIleSerLysGlyGluAsnAsnProGlyPheGlnAsnThrAspAspValGlnThrSerPh 805  
Db 2798 TATTACGAAGAGAGAAATTAATCCAGATTCCAAACACTGATGATGTTCCAGACCTCTT 2857  
QY 805 e 805  
Db 2858 T 2858  
RESULT 6  
ID AAS14880 standard; cdNA; 2920 BP.  
AC AAS14880;  
DT 20-DEC-2001 (first entry)  
XX Human cDNA encoding novel human protein NHP #1.  
DE Human; novel human protein; NHP; ss; antidiabetic; antirheumatic;  
KW antiarthritic; cytostatic; antiarteriosclerotic; vulnery;  
KW neuroprotective; nootropic; antiparkinsonian;  
KW anti-human immunodeficiency virus; antiaslathmic;  
KW hypotensive; anorectic; antinfertility; neuroleptic; anticonvulsant;  
KW antinflammatory; antipneumonia; antiparasitic; antitubercular;  
KW antineoplastic; antiproliferative; antitumor; antitoxic;  
KW immunomodulator; antiseborrheic; dermatological; vasoconstriction;  
KW gastrointestinal disorder; cardiovascular disorder; hypertension;  
KW coronary heart disease; arteriosclerosis; anorexia; obesity; bulimia;  
KW cachexia; male infertility; impotence; testicular cancer; lung tumour;  
KW hyperproliferative disorder; pulmonary system disorder;  
KW central nervous system disorder; bone disorder;  
KW neurodegenerative disease; Alzheimer's disease; Parkinson's disease;  
KW Huntington's disease; schizophrenia; mania; dementia; paranoia;  
KW panic disorder; learning disability; amyotrophic lateral sclerosis;  
KW psychosis; autism; sleep disorder; immune system disorders;  
KW Hashimoto's thyroiditis; musculo-skeletal system disorders;  
KW multiple sclerosis; ischaemic brain injury; stroke; infectious disease;  
KW diabetes mellitus; immunological disorder; asthma; AIDS;  
KW acquired immunodeficient syndrome; leukaemia; rheumatoid arthritis;

KW inflammatory bowel disease; sepsis; acne; psoriasis; lupus erythematosus;  
KW neural system disorder; respiratory disorder; olfactory disorder;  
XX wound healing; chromosome X.  
OS Homo sapiens.  
XX Key Location/Qualifiers  
FH 213..2348  
CDS  
FT /\*tag= a  
FT /product= "NHP #1"  
FT /transl\_except= (pos:867..869,aa:Xaa)  
FT /transl\_except= (pos:930..932,aa:Xaa)  
FT /transl\_except= (pos:1707..1709,aa:Xaa)  
FT /note= "Xaa= Any amino acid"  
XX WO200174896-A1.  
XX 11-OCT-2001.  
XX 02-APR-2001; 2001WO-US10542.  
XX 03-APR-2000; 2000US-194118P.  
XX 29-SEP-2000; 2000US-236384P.  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX Moore PA, Ni J, Soppet DR, Coleman TA, Gentz RL, Endress GA;  
PI Li Y, Dillon PJ;  
XX WPI: 2001-626394/72.  
DR P-PSDB; AAU09092.  
XX New human proteins, useful for diagnosing, treating, preventing and/or  
PT prognosing disorders related to the proteins, including cardiovascular  
PT disorders, autoimmune disorders and reproductive disorders -  
XX Claim 1: Page 291-292; 318pp; English.  
XX The invention relates to novel human proteins (NHP) and the  
CC nucleic acids that encode them and antibodies raised against them.  
CC The proteins, antibodies and nucleic acids are useful in the diagnosis,  
CC prognosis, prevention and/or treatment of diseases and/or disorders  
CC involving vasoconstriction, gastrointestinal disorders, cardiovascular  
CC disorders (e.g. hypertension, erectile dysfunction, high blood pressure,  
CC coronary heart disease and arteriosclerosis), anorexia, obesity, bulimia,  
CC cachexia, disorders of small intestine, disorders of reproductive system  
CC (e.g. male infertility and/or impotence), testicular cancer, lung tumours  
CC and other hyperproliferative disorders, disorders of pulmonary system,  
CC central nervous system disorders, bone disorders, neurodegenerative  
CC diseases and behavioural disorders (e.g. Alzheimer's disease, Parkinson's  
CC disease, Huntington's disease, schizophrenia, mania, dementia, paranoia,  
CC panic disorder, learning disabilities, amyotrophic lateral sclerosis,  
CC psychoses, autism, sleep disorders), immune system disorders (e.g.  
CC Hashimoto's thyroiditis), renal and musculo-skeletal system disorders,  
CC central nervous system disorders (e.g. multiple sclerosis, ischaemic  
CC brain injury and/or stroke), infectious diseases, diabetes mellitus,  
CC immunological disorders (e.g. asthma, acquired immunodeficient syndrome  
CC (AIDS), leukaemia, rheumatoid arthritis, inflammatory bowel disease,  
CC sepsis, acne, psoriasis and lupus erythematosus), neural system  
CC disorders, respiratory disorders, olfactory disorders and wound  
CC healing. The present sequence encodes an NHP of the invention and  
XX is located on the X chromosome.  
SQ Sequence 2920 BP; 897 A; 568 C; 654 G; 788 T; 13 other;  
Alignment Scores:  
Pred. No.: 0 Length: 2920  
Score: 4061.00 Matches: 763  
Percent Similarity: 99.35% Conservative: 1  
Best Local Similarity: 99.22% Mismatches: 4  
Query Match: 94.64% Indels: 1  
DB: 22 Gaps: 0



QY 722 uGluPheLeuGlyIleGlnProThrLeuGlyProProAsnGlnProProValSerIleTr 742  
 DB 2195 AGAGTTCTGGGATACAGCCACACTTGGACCTCTTACCACCCCTCTTCATATG 2254

QY 742 pLeuIleValPheGlyValValMetGlyValIleValValGlyIleValIleLeuIlePh 762  
 DB 2255 GCTGATTTGTTTGGAGTCTGATGGAGTGATAGTGGTGGCATGTCTCATCTGATCTT 2314

QY 762 eThrGlyIleArgAspArgLysLys 770  
 DB 2315 CACTGGCATCAGACATCGGAAGAAG 2339

RESULT 7  
 AAS14890  
 ID AAS14890 standard; cDNA; 2911 BP.  
 XX  
 AC AAS14890;  
 XX  
 DT 20-DEC-2001 (first entry)  
 XX  
 DE Human cDNA encoding novel human protein NHP #11.  
 XX  
 KW Human: novel human protein; NHP; ss: antidiabetic; antirheumatic;  
 KW antiarthritic; cytosolic; antiarteriosclerotic; vulnery;  
 KW neuroprotective; nootropic; antiparkinsonian;  
 KW anti-human immunodeficiency virus; antiasthmatic; vasotropic; cardiant;  
 KW hypotensive; anorectic; antinfertility; neuroleptic; anticonvulsant;  
 KW antimanic; immunosuppressive; cerebroprotective; antimicrobial;  
 KW antinflammatory; antibacterial; antipsoriatic; thymolmetic;  
 KW immunomodulator; antiseborrheic; dermatological; vasoconstriction;  
 KW gastrointestinal disorder; cardiovascular disorder; hypertension;  
 KW coronary heart disease; arteriosclerosis; anorexia; obesity; bulimia;  
 KW cachexia; male infertility; impotence; testicular cancer; lung tumour;  
 KW hyperproliferative disorder; pulmonary system disorder;  
 KW central nervous system disorder; bone disorder;  
 KW neurodegenerative disease; Alzheimer's disease; Parkinson's disease;  
 KW Huntington's disease; schizophrenia; mania; dementia; paranoia;  
 KW panic disorder; learning disability; anyotropic lateral sclerosis;  
 KW psychosis; autism; sleep disorder; immune system disorder;  
 KW Hashimoto's thyroiditis; musculo-skeletal system disorders;  
 KW multiple sclerosis; ischaemic brain injury; stroke; infectious disease;  
 KW diabetes mellitus; immunological disorder; asthma; AIDS;  
 KW acquired immunodeficient syndrome; leukaemia; rheumatoid arthritis;  
 KW inflammatory bowel disease; sepsis; acne; psoriasis; lupus erythematosus;  
 KW neural system disorder; respiratory disorder; olfactory disorder;  
 wound healing.  
 XX  
 OS Homo sapiens.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 213..998  
 FT /tag- a  
 FT /product- "NHP #11"  
 XX  
 XX WO200174896-A1.  
 XX  
 XX 11-OCT-2001.  
 XX  
 XX 02-APR-2001; 2001WO-US10542.  
 XX  
 XX 03-APR-2000; 2000US-194118P.  
 XX 29-SEP-2000; 2000US-236384P.  
 XX  
 XX (HUMA-) HUMAN GENOME SCI INC.  
 XX  
 XX Moore PA, Ni J, Soppet DR, Coleman TA, Gentz RL, Endress GA;  
 PI Li Y, Dillon PJ;  
 XX  
 XX WPI: 2001-626394/72.  
 DR P-PSDB; AAU09102.  
 XX  
 PT New human proteins, useful for diagnosing, treating, preventing and/or  
 PT prognosing disorders related to the proteins, including cardiovascular

disorders, autoimmune disorders and reproductive disorders -  
 Claim 1: Page 297-298; 318pp; English.  
 XX  
 CC The invention relates to novel human proteins (NHP) and the  
 CC nucleic acids that encode them and antibodies raised against them.  
 CC The proteins, antibodies and nucleic acids are useful in the diagnosis,  
 CC prognosis, prevention and/or treatment of diseases and/or disorders  
 CC involving vasoconstriction, gastrointestinal disorders, cardiovascular  
 CC disorders (e.g. hypertension, erectile dysfunction, high blood pressure,  
 CC coronary heart disease and arteriosclerosis), anorexia, obesity, bulimia,  
 CC cachexia, disorders of small intestine, disorders of reproductive system  
 CC (e.g. male infertility and/or impotence), testicular cancer, lung tumours  
 CC and other hyperproliferative disorders, disorders of pulmonary system,  
 CC central nervous system disorders, bone disorders, neurodegenerative  
 CC diseases and behavioural disorders (e.g. Alzheimer's disease, Parkinson's  
 CC disease, Huntington's disease, schizophrenia, mania, dementia, paranoia,  
 CC panic disorder, learning disabilities, anyotropic lateral sclerosis,  
 CC psychoses, autism, sleep disorders), immune system disorders (e.g.  
 CC Hashimoto's thyroiditis), renal and musculo-skeletal system disorders,  
 CC central nervous system disorders (e.g. multiple sclerosis, ischaemic  
 CC brain injury and/or stroke), infectious diseases, diabetes mellitus,  
 CC immunological disorders (e.g. asthma, acquired immunodeficient syndrome  
 CC (AIDS), leukaemia, rheumatoid arthritis, inflammatory bowel disease,  
 CC sepsis, acne, psoriasis and lupus erythematosus), neural system  
 CC disorders, respiratory disorders, olfactory disorders and wound  
 CC healing. The present sequence encodes an NHP of the invention.  
 XX  
 SQ Sequence 2911 BP; 896 A; 570 C; 655 G; 788 T; 2 other;  
 Alignment Scores:  
 Pred. No.: 0 Length: 2911  
 Score: 4013.00 Matches: 763  
 Percent Similarity: 99.35% Conservative: 1  
 Best Local Similarity: 99.22% Mismatches: 4  
 Query Match: 93.52% Indels: 3  
 DB: 22 Gaps: 0  
 US-09-635-501-2 (1-805) x AAS14890 (1-2911)  
 QY 3 SerSerSerTrpLeuLeuSerLeuValAlaValThrAlaAlaGlnSerThrIleGlu 22  
 DB 35 AGCTCTCTCGGCTCTCTCAGCCTTGTGCTGTAACCTGCTGCTCAGTCACCATGAG 94  
 QY 23 GluGlnAlaLysThrPheLeu-AspLysPheAsnHisGluAlaGluAspLeuPheTyrGI 42  
 DB 95 GAACAGGCCAAGACATTTTGGGACAAAGTTTAAACCAAGAGCCGACCTGTCTATCA 154  
 QY 42 nSerSerLeuAlaSerTrpAsnTrpAsnThrAsnIleThrGluGluAsnValGlnAsnMe 62  
 DB 155 AAGTTCACTTGTCTTGGGAATATTAACACCAATATTAAGAGAGATCTCCAAACAT 214  
 QY 62 tAsnAsnAlaGlyAspLysTrpSerAlaPheLeuLysGluGlnSerThrIleuAlaGlnMe 82  
 DB 215 GAATAATGCTGGGACAAATGGTCTGCTCTTTTAAAGGAACAGTCCACACTTGCCAAAT 274  
 QY 82 tTyrProLeuGlnGlnIleGlnAsnLeuThrValLysLeuGlnLeuAlaLeuGlnGI 102  
 DB 275 GTATCCACTACAAGAAATTCAGAAATCTCACAGTCAGCTTCAGCTGCGGCTCTTCAGCA 334  
 QY 102 nAsnGlySerSerValLeuSerGluAspLysSerLysArgLeuAsnThrIleLeuAsnTh 122  
 DB 335 AAATGGTCTTTCAGTCTCTCAGAGACAAAGACAAACGCTTGAACACAAATTTCAATAAC 394  
 QY 122 rMetSerThrIleTyrSerThrGlyLysValCysAsnProAspAsnProGlnGluCysLe 142  
 DB 395 AATGACACCATCTACAGTACTCGAAAAGTTTGTAAACCCAGATAATCCACGAATGCTT 454  
 QY 142 uLeuLeuGluProGlyLeuAsnGluIleMetAlaAsnSerLeuAspTyrAsnGluArgLe 162  
 DB 455 ATTACTTGACACAGGTTTGATGAATAATGGCAACAGTTTAGACTACATGAGAGGCT 514  
 QY 162 uTrpAlaTrpGluSerTrpArgSerGluValGlyLysGlnLeuArgProLeuTyrGluGI 182

Db 515 CTGGCTTGGGAAAGCTGGAGATCTGAGTGGCAAGCAGCTGAGGCCATTATATGAAGA 574  
 Qy 182 uTyrValValLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAspTyrGlyAspTy 202  
 Db 575 GTATGTGCTTGGAAATGAGATGGCAAGAGCAAAATCATTTATGAGGACTATGGGGATTA 634  
 Qy 202 rTPArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGlyGlnLe 222  
 Db 635 TTGGAGAGGAGACTATGAAGTAAATGGGGTAGATGGCTATGACTACAGCCGCGGCCAGTT 694  
 Qy 222 uileGluaspValGluHisThrPheGluGluLeuLysProLeuTyrGluHisLeuHisAl 242  
 Db 695 GATTGAAGATGTGAACATACCTTTGAAGAGATTAAACCATATATGAACATCTTCATGC 754  
 Qy 242 aTyrValArgAlaLysLeuMetAsnAlaTyrProSerTyrIleSerProIleGlyCysLe 262  
 Db 755 CTATGTGAGGCCAAAGTTGATGATGCTATCCTTCTTATATCAGTCCAAATGGATGCT 814  
 Qy 262 uProAlaHisLeuLeuGlyAspMetTyrGlyArgPheThrAsnLeuTyrSerLeuTh 282  
 Db 815 CCTGCTCATTTGCTTGTGATATCTGGGGTAGATTTTGGACAAATCTGTACTCTTTGAC 874  
 Qy 282 rValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGlnAlaTr 302  
 Db 875 AGTTCCCTTTGGACAGAAACCAACATAGATGTTACTGATGCAATGGTGGACCGCCGTG 934  
 Qy 302 pAspAlaGlnArgIlePheLysGluAlaGluLysPhePheValSerValGlyLeuProAs 322  
 Db 935 GGATGCACAGAGATATTCAAGAGGCCGCAAGATTCCT- GTATCTGTGGTCTTCCCAA 993  
 Qy 322 nMetThrGlnGlyPheThrPGLuAsnSerMetLeuThrAspProGlyAsnValGlnLysAl 342  
 Db 994 TATGACTCAAGGATTTCTGGGAAATTCATGCTAACGACCCAGGAATGTTCAAGAAAGC 1053  
 Qy 342 aValCysHisProThrAlaTrpAspLeuGlyLysGlyAspPheArgIleLeuMetCysTh 362  
 Db 1054 AGTCGTGGCATCCACAGCTTGGGACCTGGGGAAGGGCGACTTCAGGATCCTTATGTGAC 1113  
 Qy 362 rLysValThrMetAspPhePheLeuThrAlaHisHisGluMetGlyHisIleGlnTyrAs 382  
 Db 1114 AAAGTGACATGGGACGACTTCTTCACAGCTCATCATGAGATGGGCGATATCCAGTATGA 1173  
 Qy 382 pMetAlaTyrAlaAlaGlnProPheLeuLeuArgAsnGlyAlaAsnGluGlyPheHisGl 402  
 Db 1174 TATGGCATATGCTGCACAACTTTCTGCTAAGAAATGGAGCTAATGAAGGATTCATGA 1233  
 Qy 402 uAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLysHisLeuLysSerIleGl 422  
 Db 1234 AGCTGTTGGGAAATTCATGTCTACATTTCTGCAGCCACACTAAGCATTTAAATCCATTGG 1293  
 Qy 422 yLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeuLysGl 442  
 Db 1294 TCTTCTGTCACCCGATTTTCAGAGAGCAATGAACACAGATAAATCTCTGCTCAACA 1353  
 Qy 442 nAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTyrArgTrpMe 462  
 Db 1354 AGCACTACGATGTTGGGACTCTGCCATTTACTTACATGTTAGAGAAAGTGGAGGTGGAT 1413  
 Qy 462 tValPheLysGlyGluIleProLysAspClnTrpMetLysLysTyrTrpGluMetLysAr 482  
 Db 1414 GGTCTTTAAGGGGAAATTTCCCAAGACCAGTGGATGAAGAAAGTGGTGGAGATGAAGCG 1473  
 Qy 482 gGluIleValGlyValValGluProValProHisAspGluThrTyrCysAspProAlaSe 502  
 Db 1474 AGAGATAGTTGGGTGGTGGACCTGTGCCCATCATGAACATACTGTGACCCCGCATC 1533  
 Qy 502 rLeuPheHisValSerAsnAspTyrSerPheIleArgTyrThrArgThrLeuTyrGl 522  
 Db 1534 TCTGTTCATGTTCTTAATGATGATTACTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTT 1593  
 Qy 522 nPheGlnPheGlnGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHisLysCy 542

Db 1594 ATTCAGGTTTCAAGAAAGCAGCTTTGTCAAGCAGCTTAAACATGAAGCCCTCTCTGCACAAATG 1653  
 Qy 542 sasPtleSerAsnSerThrGluAlaGlyGlnLysLeuPheAsnMetLeuArgLeuGlyLy 562  
 Db 1654 TGACATCTC-AACCTACAGAGCTGGACAGAAACTGTTCAATATGCTCAGGNTTGGAAA 1712  
 Qy 562 sSerGluProTrpThrLeuAlaLeuGluAsnValValGlyAlaLysAsnMetAsnValAr 582  
 Db 1713 ATCAGAACCTGGACCTAGCATTTGGAATTTGTTAGCAGCAAGAACATCAATGTATAG 1772  
 Qy 582 gProLeuLeuAsnTyrPheGluProLeuPheThrTrpLeuLysAspGlnAsnLysAsnSe 602  
 Db 1773 GCCACTGCTCAACTACTTTGAGCCCTTATTTACCTTGGCTTGAAGACCAAGAACAGATTC 1832  
 Qy 602 rPheValGlyTrpSerThrAspTrpSerProTyrAlaAspGlnSerIleLysValArgI 622  
 Db 1833 TTTTCTGGGATGGAGTACCGACTGGAGTCCATATGCAGACCAACATCAAAAGTGGAGT 1892  
 Qy 622 eSerLeuLysSerAlaLeuGlyAspLysAlaTyrGluTrpAsnAspAsnGluMetTyrIle 642  
 Db 1893 AAGCCTAAATCAGCTCTTGGAGATAAAGCATATGAATGGAACGACAAATGAAATGTACT 1952  
 Qy 642 uPheArgSerSerValAlaTyrAlaMetArgGlnTyrPheLeuLysValLysAsnGlnMe 662  
 Db 1953 GTTCGATCATCTGTTCATATGCTATGAGGCGAGTACTTTTAAAGTAAAAATCAGAT 2012  
 Qy 662 tIleLeuPheGlyGluGluAspValArgValAlaAsnLeuLysProArgIleSerPheAs 682  
 Db 2013 GATTCCTTTTGGGAGGAGGATGCGAGTGGCTAATTTGAACCAAGAAATCTCCTTTAA 2072  
 Qy 682 nPhePheValThrAlaProLysAsnValSerAspIleIleProArgThrGluValGluLy 702  
 Db 2073 TTTCTTTGTCACCTGCACCTAAAAATGCTCTGATATCATTCCTAGAACCTGAAGTTGAAA 2132  
 Qy 702 salAlaArgMetSerArgSerArgIleAsnAspAlaPheArgLeuAsnAspAsnSerLe 722  
 Db 2133 GCCCATCAGGATGTCGGAGCGGATCATCAATGATGCTTCCGTCTGAATGACGACAGCT 2192  
 Qy 722 uGluPheLeuGlyIleGlnProThrLeuGlyProProAsnGlnProProValSerIleTr 742  
 Db 2193 AGAGTTCTTGGGGATACAGCCAACTTGGACTCTCTAACACAGCCCTGTTTCCATATG 2252  
 Qy 742 pleulleValPheGlyValValMetGlyValIleValValGlyIleValIleleuleph 762  
 Db 2253 GCTGATGTTTGGAGTGTGATGGAGTGTAGTGTGGCATTTGGCATTTGTCATCTGATCTT 2312  
 Qy 762 eThrGlyIleArgAspArgLysLys 770  
 Db 2313 CACTGGATCAGATCGGAAGAG 2337  
 RESULT 8  
 AAZ59466  
 ID AAZ59466 standard; DNA; 2262 BP.  
 XX AAZ59466;  
 AC AAZ59466;  
 DT 11-APR-2000 (first entry)  
 XX Human MPROT15 coding sequence #2.  
 XX MPROT15; treatment; hypertension; human; myocardial disease; apoplexy;  
 KW heart disease; apoplexy; heart disease; nervous denaturation; ds;  
 KW Alzheimer's disease; hormone; cytokine.  
 OS Homo sapiens.  
 XX JP11318472-A.  
 XX 24-NOV-1999.  
 XX 22-JAN-1999; 99JP-0014949.  
 XX 13-MAY-1998; 98GB-0010373.

PR 18-AUG-1996; 98GB-0018009.  
 XX (SMK ) SMITHKLINE BEECHAM PLC.  
 XX WPI; 2000-109268/10.  
 XX MPR015 polypeptide and MPR015 polynucleotides - useful for the  
 PT treatment of hypertension, myocardial diseases, apoplexy, heart  
 PT diseases, nervous denaturation, Alzheimer's disease etc.  
 XX Claim 18; Page 15; 22pp; Japanese.  
 XX This is coding sequence #2 of human MPR015. The MPR015 polynucleotide  
 CC and polypeptide sequences can be used for the treatment of hypertension,  
 CC myocardial diseases, apoplexy, heart diseases, nervous denaturation,  
 CC Alzheimer's disease and diseases related to the processing of peptide  
 CC hormones and cytokines.  
 XX SQ Sequence 2262 BP; 693 A; 450 C; 523 G; 596 T; 0 other;  
 Alignment Scores:  
 Pred. No.: 0 Length: 2262  
 Score: 3740.50 Matches: 711  
 Percent Similarity: 90.57% Conservative: 9  
 Best Local Similarity: 89.43% Mismatches: 12  
 Query Match: 87.17% Indels: 63  
 DB: 21 Gaps: 3  
 US-09-635-501-2 (1-805) x AA259466 (1-2262)  
 QY 11 LeuValAlaValThrAlaAlaGlnSerThrIleGluGluAlaLysThrPheLeuAsp 30  
 DB 64 CTTGTGCTGTAACCTGCTCAGTCCACCATTTGAGAACAGCCAGACATTTTGGAC 123  
 QY 31 LysPheAsnHisGluAlaGluAspLeuPheTyrGlnSerSerLeuAlaSerTrpAsnTyr 50  
 DB 124 AAGTTAACACGAGAGCCAGACCTGTTCTATCAAGTTCACCTGCTTCTTGGAAATAT 183  
 QY 51 AsnThrAsnIleThrGluGluAsnValGlnAsnMetAsnAlaGlyAspLysTrpSer 70  
 DB 184 AACACCAATATTACTGAGAGAGATGTCACAAACATGAATAATGCTGGGACAAATGGTCT 243  
 QY 71 AlaPheLeuLysGluGlnSerThrLeuAlaGlnMetTyrProLeuGlnGluIleGlnAsn 90  
 DB 244 GCCTTTTAAAGAACAGCTCCACACTGGCCCAAAATGTATCCACTACAGAAATTCAGAT 303  
 QY 91 LeuThrValLysLeuGlnLeuGlnAlaLeuGlnGlnAsnGlySerSerValLeuSerGlu 110  
 DB 304 CTCACAGTCAAGCTTCAGCTGAGGCTCTTCAGCAAAATGGGTCTTCAGTCTCAGAA 363  
 QY 111 AspLysSerLysArgLeuAsnThrIleLeuAsnThrMetSerThrIleTyrSerThrGly 130  
 DB 364 GACAAGAGCAACGGTTGAACCAATTCATAACATACATGAGCACCACCTACAGTACTGGA 423  
 QY 131 LysValCysAsnProAspAsnProGlnGluCysLeuLeuGluProGlyLeuAsnGlu 150  
 DB 424 AAAGTTTGPAACCCAGATATCCACAGAAATGCTATTACTTGAACACAGTTGGAATGAA 483  
 QY 151 IleMetAlaAsnSerLeuAspTyrAsnGluArgLeuTrpAlaTrpGluSerTrpArgSer 170  
 DB 484 AATAATGCAACACAGTTTACACTACATGAGAGGCTCTGGGCTTGGGAAAGCTGGAGATCT 543  
 QY 171 GluValGlyLysGlnLeuArgProLeuTyrGluGluTyrValValLeuLysAsnGluMet 190  
 DB 544 GAGGTGCGCAACGACGTGAGGCCATTTATATGAAGAGTATGTGGTCTTGAATAATGAGATG 603  
 QY 191 AlaAlaGlnAsnHisTyrGluAspTyrGlyAspTyrTrpArgGlyAspTyrGluValAsn 210  
 DB 604 GCAAGAGCAAAATCATATGAGGACTATGGGGATATTATGGAGAGGAGACATGAAGTAAAT 663  
 QY 211 GlyValAspGlyTyrAspTyrSerArgGlyGlnLeuIleGluAspValGluHisThrPhe 230  
 DB 664 GGGGTAGATGGCTATGACTACAGCCGCGCCAGTTGATTGAAGATGTGCAACATACCTTT 723

QY 231 GluGluIleLysProLeuTyrGluHisLeuHisAlaTyrValArgAlaLysLeuMetAsn 250  
 DB 724 GAAGAGATTAAACCATATATGAACATCTCATGCCTATGTGAGGCAAGATTTGATGAAT 783  
 QY 251 AlaTyrProSerTyrIleSerProIleGlyCysLeuProAlaHisLeuLeuGlyAspMet 270  
 DB 784 GCCTATCCTTCCATATATCAGTCCAAATGGATGCTCCCTGCTCATTTGTTGTTGATATG 843  
 QY 271 TrpGlyArgPheThrPheAsnLeuTyrSerLeuThrValProPheGlyGlnLysProAsn 290  
 DB 844 TGGGTAGATTTTGGACAAATCTGTACTCTTTCACAGTTCCTTTCGACAGAAACCAAC 903  
 QY 291 IleAspValThrAspAlaMetValAspGlnAlaTrpAspAlaGlnArgIlePheLysGlu 310  
 DB 904 ATAGATGTTACTGATCAATGGTGGACAGGCTGGATGCGACAGAGAAATATTTCAAGGAG 963  
 QY 311 AlaGluLysPhePheValSerValGlyLeuProAsnMetThrGlnGlyPheTrpGluAsn 330  
 DB 964 GCGGAGAGTTCTTCTATCTGTGCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 1023  
 QY 331 SerMetLeuThrAspProGlyAsnValGlnLysAlaValCysHisProThrAlaTrpAsp 350  
 DB 1024 TCCATGCTAACGAGCCAGCAATGTTGAGAAAGCACTGCTCCATCCACAGCTTGGAC 1083  
 QY 351 LeuGlyLysGlyAspPheArgIleLeuMetCysThrLysValThrMetAspAspPheLeu 370  
 DB 1084 CTGGGAAGGGGACTTCAGGATCTTATGTCACAAAGGTGACAATGGACGACTTCCTG 1143  
 QY 371 ThrAlaHisGluMetGlyHisIleGlnTyrAspMetAlaTyrAlaLagInProPhe 390  
 DB 1144 ACAGCTCATCATGATGGGCATATCCAGTATGATGATGATGATGATGATGATGATGATGATGAT 1203  
 QY 391 LeuLeuArgAsnGlyAlaAsnGluGlyPheHisGluAlaValGlyGluIleMetSerLeu 410  
 DB 1204 CTGCTAAGAAATGGACCTAATGAAGGATTCATGAAGCTGTGGGAAATCATGTCACCT 1263  
 QY 411 SerAlaAlaThrProLysHisLeuLysSerIleGlyLeuSerProAspPheGlnGlu 430  
 DB 1264 TCTGACGACACACCTAAGCATTTAAATCCATTTGGTCTTCTGTCACCCGATTTTCAAGAA 1323  
 QY 431 AspAsnGluThrGluIleAsnPheLeuLysGlnAlaLeuThrIleValGlyThrLeu 450  
 DB 1324 GACAAATGAACAGAAATTAACCTTCTGCTCAACAGCACTCAGGATGTGGGACTCTG 1383  
 QY 451 ProPheThrTyrMetLeuGluLysTrpArgTrpMetValPheLysGlyGluIleProLys 470  
 DB 1384 CCATTTACTTACATGTTAGAGAGTGGAGTGGATGCTCTTTAAAGGGGAAATTTCCAAA 1443  
 QY 471 AspGlnTrpMetLysLysTrpTrpGluMetLysArgGluIleValGlyValGluPro 490  
 DB 1444 GACCAGTGAATAAAGTGGTGGAGATGAAA----- 1476  
 QY 491 ValProHisAspGluThrTyrCysAspProAlaSerLeuPheHisValSerAsnAspTyr 510  
 DB 1476 ----- 1476  
 QY 511 SerPheIleArgTyrTyrThrArgThrLeuTyrGlnPheGlnPheGlnGluAlaLeuCys 530  
 DB 1477 -----TATTACACAGGACCTTTTACCAATTCAGTTTCAAGAAGCACTTGT 1524  
 QY 531 GlnAlaAlaLysHisGluGlyProLeuHisLysCysAspIleSerAsnSerThrGluAla 550  
 DB 1525 CAAGCAGCTTAAACATCAAGCCCTCTGCAAAATGTGACATCTCAAACTCTACAGAAGCT 1584  
 QY 551 GlyGlnLysLeuPheAsnMetLeuArgLeuGlyLysSerGluProTrpThrLeuAlaLeu 570  
 DB 1585 GGACAGAAACTGTTCAATATGCTGAGGCTTGGAAATCAAGACCTGGACCTTACGATTG 1644  
 QY 571 GluAsnValValGlyAlaLysAsnMetAsnValArgProLeuLeuAsnTyrPheGluPro 590  
 DB 1645 GAAATGTTGTTAGGAGCAAGAACATGAATGTAAAGCCACTGCTCAACTACTTTTGAGCCC 1704



QY 591 LeuPheThrTriLeuLysAspGlnAsnLysAsnSerPheValGlyTrpSerThrAspTrp 610  
Db 1705 TTATTACTGCTGCTGAAGACCAAGCAAGATCTTTGTGGATGGAGTACCGACTGG 1764  
QY 611 SerProTyrAlaAspGlnSerIleLysValArgIleSerLeuLysSerAlaLeuGlyAsp 630  
Db 1765 AGTCCATGG----- 1773  
QY 631 LysAlaTyrGluTrpAsnAspAsnGlnMetTyrLeuPheArgSerValAlaTyrAla 650  
Db 1774 -----GAAGTCTTTCATCTCTGATTGTGCTCTGTGCCA 1809  
QY 651 MetArgGlnTyrPheLeuLysValLysAsnGlnMetIleLeuPheGlyGluAspVal 670  
Db 1810 CNAAGTCAGATCTTTGTT-----TTGTTCTCTCAGGAGGAGGATGG 1854  
QY 671 ArgValAlaAsnLeuLysProArgIleSerPheAsnPhePheValThrAlaProLysAsn 690  
Db 1855 CGAGTGGCTAATTTGAACCAAGAAATCTCTTTAATTCTTGTCTACCTGACCTAAAT 1914  
QY 691 ValSerAspIleIleProArgThrGluValGluLysAlaIleArgMetSerArgSerArg 710  
Db 1915 GTGTCTGATATCATTTCTAGAACTGAAGTTGAAAGGCCATCAGGATGTCCCGAGCCGT 1974  
QY 711 IleAsnAspAlaPheArgLeuAsnAspAsnSerLeuGluPheLeuGlyIleGlnProThr 730  
Db 1975 ATCATGATGCTTTCGCTGATGACACAGCCCTAGAGTTCTCGGGATACAGCAACA 2034  
QY 731 LeuGlyProProAsnGlnProProValSerIlefrpLeuIleValPheGlyValMet 750  
Db 2035 CTTGGACCTCTTAACAGCCCTCTTTCCATATGCTGATGTTTGGAGTCTGTGATG 2094  
QY 751 GlyValIleValGlyIleValIleLeuIlePheThrGlyIleArgAspArgLysLys 770  
Db 2095 GGAGTGATAGTGGTGGCATCTCATCTGATCTTTCACGGGATCAGAGATCGGAAGAG 2154  
QY 771 LysAsnLysAlaArgSerGlyGluAsnProTyrAlaSerIleAspLysSerLysGlyGlu 790  
Db 2155 AAAAAATAAGCAAGAGTGGAGAAAAATCTTATGCTCCATCGATATTAGCAAGAGAA 2214  
QY 791 AsnAsnProGlyPheGlnAsnThrAspAspValGlnThrSerPhe 805  
Db 2215 AATAATCCAGGATCCAAACACTGATGATGTTTCAGACCTCTCTT 2259  
RESULT 9  
AAC84368  
ID AAC84368 standard; cDNA; 2638 BP.  
XX  
AC AAC84368;  
XX  
XX 19-MAR-2001 (first entry)  
XX  
DE Mouse Zace2-5 protein encoding cDNA.  
XX  
KW Zace2; metalloenzyme; angiotensin-converting enzyme; ACE; fertility;  
KW zinc metalloproteinase; blood pressure; zinc protease; hypertension;  
KW ventricular systolic dysfunction; renal impairment; heart failure;  
KW scleroderma renal crisis; atherosclerosis; antiinflammatory; mouse;  
KW antiarthritic; bradykinin inactivator; ss.  
XX  
OS Mus sp.  
XX  
FH Key  
FT CDS  
FT /tag= a  
FT /product= "Zace2-5"  
FT /note= "the coding fragment is specifically claimed for"  
XX  
PN WO200070032-A1.  
XX  
XX 23-NOV-2000.  
XX  
XX 03-MAY-2000; 2000WO-US11932.  
PF

XX 13-MAY-1999; 99US-0311482.  
PR 27-AUG-1999; 99US-0384706.  
XX (ZYMO ) ZYMOGENETICS INC.  
XX Piddington CS, Petrie CR, Shoemaker KE, Bishop PD;  
XX WPI: 2001-025018/03.  
DR P-PSDB; AAB48097.  
XX  
PT Angiotensin-converting enzyme, Zace2, useful for treating inflammatory  
PT bowel disease, e.g. Crohn's disease and ulcerative colitis, or diseases  
PT associated with inflammation such as arthritis and enterocolitis -  
XX  
XX Claim 10; Page 104-109; 125pp; English.  
PS The invention relates to the metalloenzyme Zace2. Zace2, an angiotensin-  
XX converting enzyme is a zinc metalloproteinase that plays roles in blood  
CC pressure regulation and fertility. Zace2 can be expressed by standard  
CC recombinant methodology. Zace2 polypeptides are useful for treating an  
CC inflammatory bowel disease (e.g. Crohn's disease and ulcerative colitis),  
CC diseases associated with inflammation like arthritis and enterocolitis,  
CC as targets for identifying modulators of zinc protease activity, for  
CC screening or identifying new angiotensin-converting enzyme (ACE)  
CC inhibitors, and as a basis for rational drug design for inhibitory  
CC molecules. The nucleic acids can be used to detect the expression of a  
CC Zace2 gene in a biological sample, as probes for in vivo diagnosis and  
CC for detecting and localizing Zace2 gene expression in tissue samples,  
CC to determine whether a subject's chromosomes contain a mutation in the  
CC Zace2 gene, and to detect aberrations associated with the Zace2 locus.  
CC Inhibitors of ACE are used for treating hypertension of various  
CC conditions, including left ventricular systolic dysfunction, progressive  
CC renal impairment, scleroderma renal crisis, congestive heart failure due  
CC to dysfunction, and treatment of atherosclerosis. Zace2 agonists may be  
CC used to treat infertility while Zace2 antagonists are used for inducing  
CC infertility. The present sequence represents a cDNA encoding the mouse  
CC Zace2-5 protein.  
XX  
SQ Sequence 2638 BP; 802 A; 556 C; 611 G; 669 T; 0 other;  
Alignment Scores:  
Pred. No.: 0 Length: 2638  
Score: 3579.00 Matches: 661  
Percent Similarity: 89.57% Conservative: 60  
Best Local Similarity: 82.11% Mismatches: 84  
Query Match: 83.41% Indels: 0  
DB: 22 Gaps: 0  
US-09-635-501-2 (1-805) x AAC84368 (1-2638)  
QY 1 MetSerSerSerSerTrpLeuLeuSerLeuValAlaValThrAlaGlnSerThr 20  
Db 106 ATGTCAGCTCCTCCCTGGCTCTCTCAGCTTGTGTGTACTGTCTAGTCCCTC 165  
QY 21 IleGluGluGlnAlaLysThrPheLeuAspLysPheAsnHisGluAlaGluAspLeuPhe 40  
Db 166 ACCGAGGAAAATGCCAAGACATTTTAAACAACCTTTAATCAGCAAGCTGAAGACTGTCT 225  
QY 41 TyrGlnSerSerLeuAlaSerTrpAsnTyrAsnThrAsnIleThrGluGluAsnValGln 60  
Db 226 TATCAAGTTCACTTCTCTTGGAAATTAATACTAATTAATTAATTAATTAATTAATTA 285  
QY 61 AsnMetAsnAsnAlaGlyAspLysTrpSerAlaPheLeuLysGluGlnSerThrLeuAla 80  
Db 286 AGATGATGAGCTCCAGCAATGCTGCTCTTGTGCTTTTATGAAGAACAGTCTAGACTGCC 345  
QY 81 GlnMetTyrProLeuGlnGluIleGlnAsnLeuThrValLysLeuGlnAlaLeu 100  
Db 346 CAAAGTTTCTCACTACAAGAAATCCAGACTCCGATCATCAAGCGTCAACTACAGCCCTT 405  
QY 101 GlnGlnAsnGlySerSerValLeuSerGluAspLysSerLysArgLeuAsnThrIleLeu 120  
PF



Db 406 CAGCAAGAGTGGCTCTCAGCACTCTCAGCAGACAAGAACAAACAGTTGAACACAATTCCTG 465  
Qy 121 AsnThrMetSerThrIleTyrSerThrGlyLysValIcysAsnProAspAsnProGlnGlu 140  
|||||  
Db 466 AACACGTCAGACCACTTACAGTACTGGAAGATTTGCAACCAAGAACCAACCAAGAA 525  
Qy 141 CysLeuLeuGluProGlyLeuAsnGluIleMetAlaAsnSerLeuAspTyrAsnGlu 160  
|||||  
Db 526 TGCCTATTACTTGGAGCAGGATGGATGAATAATGTGGCAGACACAGCTACAACTCT 585  
Qy 161 ArgLeuTrpAlaTrpCluserTrpArgSerGluValGlyLysGlnLeuArgProLeuTyr 180  
|||||  
Db 586 AGGCCTGGGCATGGAGGCTGGAGGCTGAGGTGGCAAGCAGCTGAGGCGCTGTGAT 645  
Qy 181 GluGluTyrValIleLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAspTyrGly 200  
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Db 646 GAAGAGTATGTGCTAGAAACAGAGATGGCAAGAGCAACAATATTAACGACATATGG 705  
Qy 201 AspTyrTrpArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGly 220  
|||||  
Db 706 GATTATTGGAGAGGAGCTATGAAGCAGAGGAGGAGCAGATGGCTACAACTATTAACCGTAAC 765  
Qy 221 GlnLeuIleGluAspValGlnHisThrPheGluGluIleLysProLeuTyrGluHisLeu 240  
|||||  
Db 766 CAGTTGATTGAAGATGTAGAAGCTACCTTCGCAGAGATCAAGCCATGTATGAGCATCT 825  
Qy 241 HisAlaTyrValArgAlaLysLeuMetAsnAlaTyrProSerTyrIleSerProIleGly 260  
|||||  
Db 826 CATGCTATGTGAGGAGGAAGTTGATGGATACCTACCTCTCTACATCAGCCCACTGGA 885  
Qy 261 CysLeuProAlaHisLeuLeuGlyAspMetTrpGlyArgPheTrpThrAsnLeuTyrSer 280  
|||||  
Db 886 TGCCTCCCTGCCATTTGCTGGTGATATGTGGGTAGATTTTGGACAATCTGTACCT 945  
Qy 281 LeuThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGln 300  
|||||  
Db 946 TTGACTGTTCCTTTGCCACAGAAACCAACATAGATGTTACTGATGCAATGATATCAG 1005  
Qy 301 AlaTrpAspAlaGlnArgIlePheLysGluAlaGluLysPhePheValSerValGlyLeu 320  
|||||  
Db 1006 GGCCTGGATGCAAGAGATATTTCAAGAGGAGAGAAATCTCTTCTCTCTGTCCTT 1065  
Qy 321 ProAsnMetThrGlnGlyPheTrpGluAsnSerMetLeuThrAspProGlyAsnValGln 340  
|||||  
Db 1066 CTTATATGACTCAAGGATCTGGCAAACTCTATGCTGACGACGACGACGATGCGCG 1125  
Qy 341 LysAlaValcysHisProThrAlaTrpAspLeuGlyLysGlyAspPheArgIleLeuMet 360  
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Db 1126 AAAGTTGTCTGCCACCCACAGCTTGGGATCTGGGACACGGAGACTTCAGAAATCAAGATG 1185  
Qy 361 CysThrLysValThrMetAspAspPheLeuThrAlaHisHisGluMetGlyHisIleGln 380  
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Db 1186 TGTACAAGGTCACATGACCACTTCTTTCACAGCCCATCAGCAGATGGGACACATCCAA 1245  
Qy 381 TyrAspMetAlaTyrAlaAlaGlnProPheLeuLeuArgAsnGlyAlaAsnGluGlyPhe 400  
|||||  
Db 1246 TATGACATGGCATATGCCAGCAACCTTCTCTGCTAAGAAACGGAGCCCAATGAAGGCTC 1305  
Qy 401 HisGluAlaValGlyLysLeuMetSerLeuSerAlaAlaThrProLysHisLeuLysSer 420  
|||||  
Db 1306 CATGAAGCTGTGGAGAAATCATGTCTACCTTCTGACGCTACCCCAAGCATCTGAAATCC 1365  
Qy 421 IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluLeuAsnPheLeuLeu 440  
|||||  
Db 1366 ATGGTCTCTGCCATTCGGATTTCAAGAAAGATAGGAAACAGAGATAACTTCTTACTG 1425  
Qy 441 LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTrpArg 460  
|||||  
Db 1426 AAACAGGCATTCACAATTTCTGGAACACTACCTTTTACTATCATGTATTAGAAAGTGGAGG 1485  
Qy 461 TrpMetValPheLysGlyLysLeuProLysAspGlnTrpMetLysLysTrpTrpGluMet 480  
|||||  
Db 1486 TGGATGGCTTTTCCGGGGTGAATTCCTCAAGAGCAGCTGGATGAAAGAGTGGTGGGAGATG 1545

RESULT 10  
AAC84370  
ID AAC84370 standard: cDNA; 2638 BP.  
XX

Qy 481 LysArgGluIleValGlyValGluProValProHisAspGluThrTyrCysAspPro 500  
|||||  
Db 1546 AAGCGGGAGATCGTTGGTGGTGGAGCCTCTGCCTCATGATGAACATCTGTGACCCT 1605  
Qy 501 AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrTyrThrArgThrLeu 520  
|||||  
Db 1606 GCATCTCTGTCCATGTTTCTAATGATTACTCATTCATTCATGATATTACACAGGACCAT 1665  
Qy 521 TyrGlnPheGluPheGlnGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis 540  
|||||  
Db 1666 TACCAATTCCAGTTTCAAGAGCTCTTGTCAAGCAGCTAAGTATATGTTCTCTCTGCAC 1725  
Qy 541 LysCysAspIleSerAsnSerThrGluAlaGlyGlnLysLeuPheAsnMetLeuArgLeu 560  
|||||  
Db 1726 AAATGTGACATCTCAATTCCTCAAGCTGGGAGAGTTGCTCAAGATGCTGAGTCTT 1785  
Qy 561 GlyLysSerGluProTrpThrLeuAlaLeuGluAsnValValGlyAlaLysAsnMetAsn 580  
|||||  
Db 1786 GGAAATTCAGAGCCCTGGACCAAGCCTTGGAAATGTGGTAGGACCAAGGAATATGGAT 1845  
Qy 581 ValArgProLeuLeuAsnTyrPheGluProLeuPheThrTrpLeuLysAspGlnAsnLys 600  
|||||  
Db 1846 GTAAACCACTGCTCAATTACTTCCAAACCGTTGTTGACTGGCTGAAGAGCAGACAGA 1905  
Qy 601 AsnSerPheValGlyTrpSerThrAspTrpSerProTyrAlaAspGlnSerIleLysVal 620  
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Db 1906 AATTCCTTTGTGGGTGGAACACTGAATGGAGCCCATATGCCGACCAAGCATTAAGTG 1965  
Qy 621 ArgIleSerLeuLysSerAlaLeuGlyAspLysAlaTyrGluTrpAsnAspAsnGluMet 640  
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Db 1966 AGATTAAGCCTTAATACAGCTCTGGAGCTAATGCATATGAATGGACCAACCAAGCAANTG 2025  
Qy 641 TyrLeuPheArgSerSerValAlaTyrAlaMetArgGlnTyrPheLeuLysValLysAsn 660  
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Db 2026 TTCTCTGTCGATCATCTGTGCATATGCCATGAGAAGTATTTTCAATAATCAAAAAC 2085  
Qy 661 GlnMetIleLeuPheGlyGluGluAspValArgValAlaAsnLeuLysProArgIleSer 680  
|||||  
Db 2086 CAGACAGTCTCTTTCTAGAGGAAGATGACGAGTGGCGATTTGAACCAACAGAGTCTCC 2145  
Qy 681 PheAsnPhePheValThrAlaProLysAsnValSerAspIleIleProArgThrGluVal 700  
|||||  
Db 2146 TTCTACTTCTTGTCACTCACCCCAAAATGTCTGTGATGTCATTCTTAGAAGTGAAGTT 2205  
Qy 701 GluLysAlaIleArgMetSerArgSerArgIleAsnAspAlaPheArgLeuAsnAspAsn 720  
|||||  
Db 2206 GAAGATGCCATCAGGATGCTCGGGCCGCATCAATGATGTCCTTTGGCCTGAATGATAAC 2265  
Qy 721 SerLeuGluPheLeuGlyIleGlnProThrLeuGlyProProAsnGlnProProValSer 740  
|||||  
Db 2266 AGCCTGGAGTTTCTGGGGATTCACCAACACACTTGAGCCACCTTACCACCTCTCTGTACC 2325  
Qy 741 IleTrpLeuIleValPheGlyValValMetGlyValIleValValGlyIleValIleLeu 760  
|||||  
Db 2326 ATATGCTGATATTATTGTTGTTGTGTGATGGCCTGCTAGTGTGCTGCTCATCATCTG 2385  
Qy 761 IlePheThrGlyIleArgAspArgLysLysAsnLysAlaArgSerGlyGluLeuAsnPro 780  
|||||  
Db 2386 ATTGTCACTGGGATCAAGGTCGAAAGAGAAATAATGAACAAAAAGAGAGAACCCCT 2445  
Qy 781 TyrAlaSerIleAspIleSerLysGlyGluAsnAsnProGlyPheGlnAsnThrAspAsp 800  
|||||  
Db 2446 TATGACTCGATGGACATTTGGAAAGAGAGAAAGCAATGCAGGATTCCAAAACATGATGAT 2505  
Qy 801 ValGlnThrSerPhe 805  
|||||  
Db 2506 GCTCAGACTTCCTTT 2520



QY 361 CysThrLysValThrMetAspPheLeuThrAlaHisHisGluMetGlyHisIleGln 380  
Db TGTACAAAGGTGCAATGGACAACTTCTTGACAGCCCATCAGAGATGGGACACATCCAA 1245  
QY 381 TyrAspMetAlaTyrAlaAlaGlnProPheLeuArgAsnGlyValaAsnGluGlyPhe 400  
Db TATGACATGCATATGCCAGCAACCTTCTCTGCTAAAGAAAGGAGCCCAATGAAGGTTTC 1305  
QY 401 HisGluAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLysHisLeuLysSer 420  
Db CATGAAGCTGTGGAGAAATCATGTCACTTCTGAGCTACCCCAAGCATCTGAATACC 1365  
QY 421 IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeuLeu 440  
Db ATTTGGTCTTCTGCCATCCGATTTTCAAGAAGATAGCGAAACAGAGATAAACTTCCTACTG 1425  
QY 441 LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTyrArg 460  
Db AAACAGGCATTTGACAATTTGGAACACTACCGTTTACTTACATGTTAGAGAAAGTGAGG 1485  
QY 461 TrpMetValPheLysGlyGluIleProLysAspGlnTrpMetLysLysTyrTrpGluMet 480  
Db TGGATGGTCTTTCGGGGTGAATTTCCCAAGAGCAGTGGATGAAAAGTGGTGGAGATG 1545  
QY 481 LysArgGluIleValGlyValValGluProValProHisAspGluThrTyrCysAspPro 500  
Db AAGCGGAGATCCTTGGTGTGGAGCCTCTCGCTCGATGAACATACATCTGTGACCCT 1605  
QY 501 AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrTrpThrArgThrLeu 520  
Db GCATCTCTGTTCCTGTTCTTAATGATTAATCTCATTCATTCATTCATTCATTCATTCAT 1665  
QY 521 TyrGlnPheGlnPheGlnGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis 540  
Db TACCAATTCACAGTTCACAGAGCTTTTGTCAAGCAGCTTAAGTATTAATGTTCTCTGCAC 1725  
QY 541 LysCysAspIleSerAsnSerThrGluAlaGlyGlnLysLeuPheAsnMetLeuArgLeu 560  
Db AAATGTGACATCTCAATTCCTCACTGAGCTGGCAGAGTTGCTCAAGATGCTGAGTCTT 1785  
QY 561 GlyLysSerGluProTrpThrLeuAlaLeuGluAsnValValGlyAlaLysAsnMetAsn 580  
Db GGAAATTCAGAGCCCTGGACCGGCTTGGAAATGTGGTAGGAGCAAGAAATATGGAT 1845  
QY 581 ValArgProLeuLeuAsnTyrPheGluProLeuPheThrTrpLeuLysAspGlnAsnLys 600  
Db GTAAACCACTGCTCAATTTACTTCCAAACCGTTTGTGACCTGGCTGAAAGAGCAGAACAGA 1905  
QY 601 AsnSerPheValGlyTrpSerThrAspTrpSerProTyrAlaAspGlnSerIleLysVal 620  
Db AATTTCTTTTGGGGTGGAACTGTAATGGAGCCCATATGCCGACCAAGCATTTAAAGTG 1965  
QY 621 ArgIleSerLeuLysSerAlaLeuGlyAspLysAlaTyrGluTrpAsnAspAsnGluMet 640  
Db AGGATAAGCCTAAATACAGCTCTTGGAGCTAATGCATATGAATGGACCAACAGAAATG 2025  
QY 641 TyrLeuPheArgSerSerValAlaTyrAlaMetArgGlnTyrPheLeuLysValLysAsn 660  
Db TTCTCTGTTCCGATCTCTGTGCATATGCCATGAGAAAGTATCTTCAATATCAAAAAC 2085  
QY 661 GlnMetIleLeuPheGlyGluGluAspValArgValAlaAsnLeuLysProArgIleSer 680  
Db CAGACAGTTCCTTTCTAGAGGAGATGTACGAGTGAGTGAATTTGAAACCAAGAGTCTCC 2145  
QY 681 PheAsnPhePheValThrAlaProLysAsnValSerAspIleIleProArgThrGluVal 700  
Db TTCTACTCTTGTTCACCTCACCCCAAAATGTGCTGTGATCTCTCTAGAGTCAAGT 2205  
QY 701 GluLysAlaIleArgMetSerArgSerArgIleAsnAspAlaPheArgLeuAsnAspAsn 720  
Db GAAGATGCCATCAGGATGTCTCGGGCCGCATCAATGATGTCTTTGGCCTGAATGATAAC 2265

QY 721 SerLeuGluPheLeuGlyIleGlnProThrLeuGlyProProAsnGlnProProValSer 740  
Db AGCTGGAGTTCTGGGATTCAACCAACACTTGAGCCACTTACCAGCCTCCTGTCAAC 2325  
QY 741 IleTrpLeuIleValPheGlyValValMetGlyValIleValValGlyIleValIleLeu 760  
Db ATATGGCTGATATTTTGGTGTGTGTGATGGCCTGCTAGTGGTGGTGCATCATCTG 2385  
QY 761 IlePheThrGlyIleArgAspArgLysLysLysLysAlaArgSerGlyGluAsnPro 780  
Db ATTTGCACTGGGATCAACGGTCGAAAGAGAAATAATGAAACAAAGAGAGAGACCT 2445  
QY 781 TyrAlaSerIleAspIleSerLysGlyGluAsnAsnProGlyPheGlnAsnThrAspAsp 800  
Db TATGACTCGATCGACATTCGAAAGAGGAAAGCAATCAGGATTCGAAACAGTATGAT 2505  
QY 801 ValGlnThrSerPhe 805  
Db GCTCAGACTTCCTTT 2520  
RESULT 11  
AAC84367  
ID AAC84367 standard; DNA; 2415 BP.  
XX AAC84367;  
AC AAC84367;  
XX 19-MAR-2001 (first entry)  
DE Human Zace2 protein encoding degenerate sequence.  
XX Zace2; metalloenzyme; angiotensin-converting enzyme; ACE; fertility;  
KW zinc metalloproteinase; blood pressure; zinc protease; hypertension;  
KW ventricular systolic dysfunction; renal impairment; heart failure;  
KW scleroderma renal crisis; atherosclerosis; . antiinflammatory; human;  
KW antiarthritic; bradykinin inactivator; ds.  
XX Homo sapiens.  
OS WO200070032-A1.  
PN 23-NOV-2000.  
PD 03-MAY-2000; 2000WO-US11932.  
PF 13-MAY-1999; 99US-0311482.  
PR 27-AUG-1999; 99US-0384706.  
XX (ZYMO ) ZYMOGENETICS INC.  
XX Piddington CS, Petrle CR, Shoemaker KE, Bishop PD;  
PI WPI; 2001-025018/03.  
XX P-PSDB; AAB48095.  
XX Angiotensin-converting enzyme, Zace2, useful for treating inflammatory  
PT bowel disease, e.g. Crohn's disease and ulcerative colitis, or diseases  
PT associated with inflammation such as arthritis and enterocolitis -  
XX Disclosure; Page 103-104; 125pp; English.  
XX The invention relates to the metalloenzyme Zace2. Zace2, an angiotensin-  
CC converting enzyme is a zinc metalloproteinase that plays roles in blood  
CC pressure regulation and fertility. Zace2 can be expressed by standard  
CC recombinant methodology. Zace2 polypeptides are useful for treating an  
CC inflammatory bowel disease (e.g. Crohn's disease and ulcerative colitis),  
CC diseases associated with inflammation like arthritis and enterocolitis,  
CC as targets for identifying modulators of zinc protease activity, for  
CC screening or identifying new angiotensin-converting enzyme (ACE)  
CC inhibitors, and as a basis for rational drug design for inhibitory  
CC molecules. The nucleic acids can be used to detect the expression of a  
CC Zace2 gene in a biological sample, as probes for in vivo diagnosis and  
CC for detecting and localizing Zace2 gene expression in tissue samples,  
CC to determine whether a subject's chromosomes contain a mutation in the

CC Zace2 gene, and to detect aberrations associated with the Zace2 locus.  
CC Inhibitors of ACE are used for treating hypertension of various  
CC conditions, including left ventricular systolic dysfunction, progressive  
CC renal impairment, scleroderma renal crisis, congestive heart failure due  
CC to dysfunction, and treatment of atherosclerosis. Zace2 agonists may be  
CC used to treat infertility while Zace2 antagonists are used for inducing  
CC infertility. The present sequence represents a degenerate sequence  
CC encoding the human Zace2 protein.  
XX  
SQ Sequence 2415 BP; 494 A; 218 C; 398 G; 335 T; 970 other;

Alignment Scores:

Pred. No.: 0 Length: 2415  
Score: 3509.00 Matches: 644  
Percent Similarity: 80.00% Conservatives: 0  
Best Local Similarity: 80.00% Mismatches: 161  
Query Match: 81.78% Indels: 0  
DB: 22 Gaps: 0

US-09-635-501-2 (1-805) x AAC84367 (1-2415)

QY 1 MetSerSerSerThrLeuLeuLeuSerLeuValAlaValThrAlaAlaGlnSerThr 20  
DB 1 ATGWSNWSNWSNTGGVNTYNTNWSNYTNGTNGCNGTNCNGCNGCNCARWSNACN 60  
QY 21 IleGluGluGlnAlaLysThrPheLeuAspLysPheAsnHisGluAlaGluAspLeuPhe 40  
DB 61 ATHGARGARGCARGCNAARACNTTYTNGAYAAATTTAAAYCAYGARGCNGARGAYTNTY 120  
QY 41 TyrGlnSerSerLeuAlaSerTrpAsnTyrAsnThrAsnIleThrGluGluAsnValGln 60  
DB 121 TAYCARWSNWSNTGWSNWSNTGGAAATYATAYACNAAYATHACNGARGAAYGTNCAR 180  
QY 61 AsnMetAsnAlaGlyAspLysTrpSerAlaPheLeuLysGluGlnSerThrLeuAla 80  
DB 181 AAYATGAAYAGCGNGGNGAYAAATGGWSNGCNTTYTNAARGARCARWSNACNTNGCN 240  
QY 81 GlnMetYrProLeuGlnGluIleGlnAsnLeuThrValLysLeuGlnAlaLeu 100  
DB 241 CARATGTAYCCNTYCARGARATHCARAAAYTNCNGTNAARYTNCARYTNCARGCNTN 300  
QY 101 GlnGlnAsnGlySerSerValLeuSerGluAspLysSerLysArgLeuAsnThrIleLeu 120  
DB 301 CARCARAAYGGWSNWSNTNTNWSNGARGAAYARWSNAARMGNTTAAAYACNATHYN 360  
QY 121 AsnThrMetSerThrIleTyrSerThrGlyLysValCysAsnProAspAsnProGlnGlu 140  
DB 361 AAYACNATGWSNACNATHATYWSNACNGNNAARGTNTGYAAYCCNGAYAACNCARGAR 420  
QY 141 CysLeuLeuLeuGluProGlyLeuAsnGluIleMetAlaAsnSerLeuAspTyrAsnGlu 160  
DB 421 TGYTNTYNTNGARCCNGGNTNAAAYGARATHATGCGNAAYSNNTYNGAYTAYAAAYGAR 480  
QY 161 ArgLeuTrpAlaTrpGluSerTrpArgSerGluValGlyLysGlnLeuArgProLeuTyr 180  
DB 481 MGNTYNTGCGCCTGGGARGWSNTGGGWSNGARGTNGGNNAARCARNTNMGCNTNTAY 540  
QY 181 GluGluTyrValValLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAspTyrGly 200  
DB 541 GARGARTAYGTNGTNTAARAAYGARATGCGCNGCNGCNAAYCAYTAYGARGAYTAYGNG 600  
QY 201 AspTyrTrpArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGly 220  
DB 601 GAYTAYTGGMGNGNGAYTAYGARGTNAAYGGNGTNGAYGNGTAYGAYTAYWSNMGNGN 660  
QY 221 GlnLeuIleGluAspValGluHisThrPheGluGluIleLysProLeuTyrGluHisLeu 240  
DB 661 CARNTNATHGARGAYGTNGARCAACNTTYTGARGARATHAARCCNTNTAYGARCAYTNT 720  
QY 241 HisAlaTyrValArgAlaLysLeuMetAsnAlaTyrProSerTyrIleSerProIleGly 260  
DB 721 CAYGNTAYGTNMGNGCNAARYTNTATGAAYGCNTATCYCCNWSNTAYATWSNCCNATHGNG 780

QY 261 CysLeuProAlaHisLeuLeuGlyAspMetTrpGlyArgPheTrpThrAsnLeuTyrSer 280  
DB 781 TGYTNTCCNCNCAYTNTYNTNGGNGAYATGCGGCGMGNTTYTGGACNAAYTNTAYWSN 840  
QY 281 LeuThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGln 300  
DB 841 YTNACNGTNCNTTYGCGCARAARCCNAAAYATHGAYGTNACNGAYGCNATGTTNGAYCAR 900  
QY 301 AlaTrpAspAlaGlnArgIlePheLysGluAlaGluLysPhePheValSerValGlyLeu 320  
DB 901 GCNTGGGAYGCNCARMGNATHTTAAARGCNGCARAARTTYTGTNNWSNGTNGGNTYN 960  
QY 321 ProAsnMetThrGlnGlyPheTrpGluAsnSerMetLeuThrAspProGlyAsnValGln 340  
DB 961 CCNAAATGACNARGGNTTYTGGARAAAYWSNATGYTNACNGAYCCNGGNAAYGTNCAR 1020  
QY 341 LysAlaValCysHisProThrAlaTrpAspLeuGlyLysGlyAspPheArgIleLeuMet 360  
DB 1021 AARGCNGTNTGYCAYCCNACNGCNGTGGGAYTNGGNAARGGNGAYTTCGNATHYTNATG 1080  
QY 361 CysThrLysValThrMetAspPheLeuThrAlaHisHisGluMetGlyHisIleGln 380  
DB 1081 TGYACNAAARGTNCACNATGGGAYGVTYTNACNGCNCAYCAYGARATGGGNCAYATHCAR 1140  
QY 381 TyrAspMetAlaTyrAlaAlaGlnProPheLeuLeuArgAsnGlyAlaAsnGluGlyPhe 400  
DB 1141 TAYGATATGCGNTAYGCGNCARCCNTTYTNTYTMGNAAYGGGNCNAAYGARGGNTTY 1200  
QY 401 HisGluAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLysHisLeuLysSer 420  
DB 1201 CAYGARGCNGTNGGARGARATHATGWSNTNWSNGCNGCNCNAAARCAAYTNAARWSN 1260  
QY 421 IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeuLeu 440  
DB 1261 ATGGGNTYNTNWSNCCNGAYTTCARGARGAAYAGARACNGARATHAAYTNTYNTYN 1320  
QY 441 LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTrpArg 460  
DB 1321 AARCARCNCNTNACNATHGTNGGNACNTNCCNTTYACNTATYATGYTNGARAARTGCMGN 1380  
QY 461 TrpMetValPheLysGlyGluIleProLysAspGlnTrpMetLysLysTrpTrpGluMet 480  
DB 1381 TGGATGTTNTYAAAGGNGARGATHCCNNAARGAYCARTGATGAARAARTGGTGGGARATG 1440  
QY 481 LysArgGluIleValGlyValValGluProValProHisAspGluThrTyrCysAspPro 500  
DB 1441 AARMGNGARATHGTNGGNGTNGTNGARCCNTNCCNCAYGARGACNTAYTGYGAYCCN 1500  
QY 501 AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrTyrThrArgThrLeu 520  
DB 1501 GCNWSNTNTTYCAYGTNWSNAAYGAYTAYWSNTTYATHMGNTAYTAYACNMGNACNTYN 1560  
QY 521 TyrGlnPheGlnPheGlnGluAlaLeuCysGlnAlaAlaLysHisGlyProLeuHis 540  
DB 1561 TAYCARTTYCARTTYCARGARGCNTYNTGYCARGCNGCNCARCAAYCARGCNCNTNCAY 1620  
QY 541 LysCysAspIleSerAsnSerThrGluAlaGlyGlnLysLeuPheAsnMetLeuArgLeu 560  
DB 1621 AARTGYCAYATHWSNAAAYWSNACNGCNGCNGCARAARYTNTTAAATGYTNGMNTYN 1680  
QY 561 GlyLysSerGluProTrpThrLeuAlaLeuGluAsnValValClyAlaLysAsnMetAsn 580  
DB 1681 GGNAAARWSNARGCNCCTGGACNTYNGACNTNGARAAAYGTNGTNGGNGCNAARAAYATGAAY 1740  
QY 581 ValArgProLeuLeuAsnTyrPheGluProLeuPheThrTrpLeuLysAspGlnAsnLys 600  
DB 1741 GTNMGNCNTNTYNTNAAAYTAYTTCARGCNCNTTNTYACNTGGTNTAARGAYCARAAAYAR 1800  
QY 601 AsnSerPheValGlyTrpSerThrAspTrpSerProTyrAlaAspGlnSerIleLysVal 620  
DB 1801 AATWSNTYGTNGTNGWSNACNGAYTGGWSNCCNTATCGCNGAYCARWSNATHAARGTN 1860  
QY 621 ArgIleSerLeuLysSerAlaLeuGlyAspLysAlaTyrGluTrpAsnAspAsnGluMet 640



Db 453 AGCCACCAATCCTAACAGTAAGTGGAAACGTCGTAAACCCAGATAATCCACAAGAATGCT 512  
QY 142 euLeuLeuGluProGlyLeuAsnGluIleMetAlaAsnSerLeuAspTyrAsnGluArgL 162  
Db 513 TATTACTTGAACCGTTTGAATGAATAATGGCAACAGTTTAGACTACAATGAGAGGC 572  
QY 162 euTIPAlaTrpGluSerTrpArgSerGluValGlyLysGlnLeuArgProLeuTyrGluG 182  
Db 573 TCTGGCTTGGAAAGCTGGAGACTGAGGTGGCAAGAGCTGAGGCCATTATAGAAG 632  
QY 182 luTyrValValLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAsp-Tyr-GlyAs 201  
Db 633 AGTATGTGCTTGAATAATGAGTGGCAAGAGCAATCATATTAGAGACTTATTGGGA 692  
QY 201 pTyrTrpArgGlyAspTyrGluValAsnGlyValAsp---GlyTyrAspTyrSerArgL 220  
Db 693 TTATTGGAGAGAGACTATGAAGTAATGGGTAAATAGTGGATATGATTACAGCCGG 752  
QY 220 yGlnLeuIleGluAspValGluHisThr-PheGluGluIleLysProLeuTyr-GluHis 239  
Db 753 CCAGTTGATGAAGATGTGGAACATACCTGTTGAAGAGATTAAACCATTTGATAGGAACAT 812  
QY 240 LeuHisAlaTyrValArgAlaLysLeuMetAsnAlaTyrProSerTyrIleSerProIle 259  
Db 813 CTTCAGCCCTATGTGAGGCCAAGTTGATGATGCTATCCTTCCTATATCAGTCCAATT 872  
QY 260 GlyCysLeuProAlaHisLeuLeuGlyAspMetTrp-GlyArgPheTrpThrAsnLeuTy 279  
Db 873 GGATGCTCCTGCTCAATTGCTTGGTATATGTCGGGTAGATTTTGGACAATCTGTA 932  
QY 279 rSerLeuThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValas 299  
Db 933 CTCTTTGACAGTCTCCTTTGGACAGAACAAACATAGATGTTACTGATGCAATGGTGA 992  
QY 299 pGlnAlaTrpAspAlaGlnArgIlePheLysGluAla-GluLysPhePheValSerValG 319  
Db 993 CCAGCCCTGGGATGCACAGAGAATATTCAAGGAGTCCGACAGAACTCTTTGTATCTGTTG 1052  
QY 319 lyLeuProAsnMetThrGlnGlyPheTrp-GluAsnSerMetLeu-ThrAspProGlyAs 338  
Db 1053 GTCTTCCTTATGACTCTAGATTCTGCGCGCAAAATCCATGCTATACGAGCCACAGAAA 1112  
QY 338 nValGlnLysAlaValCysHis-ProThrAlaTrpAspLeuGlyLysGlyAspPheArg- 357  
Db 1113 TGTTCAGAAAGCACTCTGCCATCCCAACAGCTTGGACCTGGGAGGGCGACTTCAGAG 1172  
QY 358 IleLeuMetCysThrLys-ValThrMetAspPheLeuThrAlaHisGluMetG 377  
Db 1173 ATCCTTATGTGCACAAAGGGTAACAATGGACGACTCTCTGACAGCTCATCATGAGATGGG 1232  
QY 377 yHisIleGlnTyrAspMetAlaTyrAlaAlaGlnPro-PheLeuLeuArg-AsnGlyAla 396  
Db 1233 GCATATCCAGTATGATATGGCATATGCGCGCAACCTTTTCTGCTAAGGAAATGGAGCT 1292  
QY 397 -AsnGluGlyPheHisGluAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLy 416  
Db 1293 TAATGAAGATTCCATGAGCTGTGGGAAATCATGTCTCTGCGACCCACACTAA 1352  
QY 416 sHisLeuLysSerIleGlyLeuLeuSerProAspPheGln---GluAspAsnGluThrG 435  
Db 1353 GCATTTAAATCCATTGCTCTCTGTCCACCGAGTTTCAACGAGCAACATCAACACAGA 1412  
QY 435 uIleAsnPheLeuLysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMe 455  
Db 1413 AATAAACCTTCTGCTCAACAGCACTCACGATTGTTGGGACTTGCCATTTACTTACAT 1472  
QY 455 tLeuGluLysTrpArgTrpMetValPheLys-GlyGluIleProLysAspGlnTrpMetL 475  
Db 1473 GTTAGAAGTGGAGGTGGTGGTCTTTAAACGGGGAAATTCCTCAAGAGCCAGCTGGGTGA 1532  
QY 475 ysLys-TrpTrpGluMetLysArgGlu-IleValGlyVal-ValGluProValProHis 494  
Db 1533 AAAAGGTGGTGGAGATGAACGGAAGAATAGTTGGGGTGTGGAACTGTGCCCATG 1592

QY 494 spGluThrTyr-CysAspProAlaSerLeuPheHisValSerAsnAspTyrSerPheIle 513  
Db 1593 ATGAACATATCTGTGACCCCGCATCTCTGTTCCATCTGTTCTAATGATTACTCATTCATT 1652  
QY 514 ArgTyrTyrThrArgThrLeu-TyrGlnPheGlnPheGln-GluAlaLeu-CysGlnAla 532  
Db 1653 CGATATTACCAAGAGGCCCTGTTACCAATTCAGTTTCAAAAGAACGACTTTTGTCAAGCA 1712  
QY 533 AlaLysHisGluGlyProLeuHisLys-CysAspIle-SerAsnSerThrGlu---AlaG 551  
Db 1713 GCTAAACATGAGGCCCTCTGCACAAATTTGACATTTCAAAATTTACAGAAGCTCGTG 1772  
QY 551 lyGlnLys-LeuPheAsnMetLeuArgLeuGlyLys-SerGluProTrpThrLeuAlaLe 570  
Db 1773 GACAGAACACTGTTCAATATGCTGAGGCTTGGAAAACCTCAGAACCTGGACCCCTAGCAT 1832  
QY 570 uGluAsnValVal-GlyAlaLysAsnMetAsnValArgPro-LeuLeuAsnTyrPheGlu 589  
Db 1833 GGAATATGTTTGAAGGACCAAGAACATGAATGTAAAGCCACTGCTCAACTACTTTGAG 1892  
QY 590 ProLeuPheThrTrpLeuLysAspGlnAsnLysAsnSerPheValGlyTrpSerThrAsp 609  
Db 1893 CCTTATTTACCTGGCTGGAAGACCAAGAAATCTTTTGTGGGATGGAGTACCGAC 1952  
QY 610 TrpSerProTyrAlaAspGlnSerIle-LysValArgIleSerLeuLysSerAlaLeuG 629  
Db 1953 TGGAGTCCATATGACAGACCACAGCATCAAGTGGAGATAAGCCTAATAATCAGCTCTTG 2012  
QY 629 y-AspLysAlaTyrGluTrpAsnAsp-AsnGluMetTyrLeuPheArgSerSer-ValAl 648  
Db 2013 CAGATAAGCATATGAATGGAACGCCAATCAATGTACCTCTTCGATCACTGTTGGTGG 2072  
QY 648 aTyrAla---MetArgGlnTyrPheLeu-LysValLysAsnGlnMetIleLeuPheGlyG 667  
Db 2073 ATATTCTTAATTCAGGCAGTACTTTTAAACAAGTAAAAATCAGATGATTCTTTTGGGG 2132  
QY 667 luGluAspValArgValAlaAsnLeuLysProArgIleSerPheAsnPhePheValThrA 687  
Db 2133 AGGAGATGTGCGAGTGGCTAATTTGAACCAAGAAATCTCCTTTAATTTCTTTGTCACTG 2192  
QY 687 laProLysAsnValSer-AspIleIleProArg-ThrGluValGluLysAlaIleArgMe 706  
Db 2193 CACCTAAAATGTGCTGGATATATCTCTAGAAACTCAAGTTGAAAGGCCATCAGGAT 2252  
QY 706 tSerArgSerArg-IleAsnAspAlaPheArgLeuAsnAspAsnSerLeuGluPheLeuG 726  
Db 2253 GTCCCGAGCGTACTCCATGATGCTTTCCGTCCTGAATGACGACAGCCTAGAGTTCTG 2312  
QY 726 lyIleGlnProThrLeuGlyProAsnGlnProProValSerIleTrpLeuIleValP 746  
Db 2313 GGATACACCCACACTTGGACCTCTAACCCAGCCCTGTTTCCATATGGCTGATTGTTT 2372  
QY 746 heGlyValValMetGlyValIleValValGly-IleValIleLeu-IlePheThrGlyI 765  
Db 2373 TTGGAGTTGTGATGGGAGTGATAATTTTGGCCATGGTCTCCTGGATTTCCACTGGAAT 2432  
QY 765 eArgAspArgLysLysAsnLysAlaArgSerGlyGlu-AsnPro-TyrAlaSerIle 784  
Db 2433 CAGAGNTCCGAGAAGAAAATAAAGCAAGAGTGGAGAATAATCCTTTATGCCCTCCATC 2492  
QY 785 AspIleSerLysGlyGlu---AsnAsnProGlyPheGlnAsnThrAspAspValGlnThrS 804  
Db 2493 GATATTAGTAAGGAGTATAATAATCCAGGATTCGAGGATCCGAAACACTGATGATGTCAGACCT 2552  
QY 804 erPhe 805  
Db 2553 CCTTT 2557  
RESULT 13  
AAC84369  
ID AAC84369 standard; DNA; 2415 BP.  
XX







Qy	381	TyrAspMetAlaTyrTrpAlaAlaGlnProPheLeuLeuArgAsnGlyAlaAsnGluGlyPhe	400
Db	1141	TAYGAYATGCNTAYCGNMGNCARCCNTTYTYTNTNGNAAAYGGGNCNAAAYGARGGNTTY	1200
Qy	401	HisGluAlaValGlyGluIleMetSerLeuSerAlaAlaThrProLysHisLeuLysSer	420
Db	1201	CAYGARGCNTGGNGGARATHATGWSNYTNWSGNGCNCNACNCNAAARCAYYTNAARWSN	1260
Qy	421	IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeuLeu	440
Db	1261	ATHGGNYTNTCCNWSNGAYTTYCARGAGAYWSNCARACNACGARATHAAYTTYTYNTN	1320
Qy	441	LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTrpArg	460
Db	1321	AARCAARGCNTYACNATHGTTNGNACNYTNCCTTYACNTAYATGYTNGARAARTGCMGN	1380
Qy	461	TrpMetValPheLysGlyGluIleProLysAspGlnTrpMetLysLysTrpTrpGluMet	480
Db	1381	TGCATGGTNTTYMGNGGNGARATHCCNAAARGARCARTGGATGAARARTGCTGGGARATG	1440
Qy	481	LysArgGluIleValGlyValValGluProValProHisAspGluThrTyrCysAspPro	500
Db	1441	AARMGNGARATHGTTGGNGTNGTGARCCNYTNCNCAYGAYGARACNTAYTCYGAYCCN	1500
Qy	501	AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrTrpThrArgThrLeu	520
Db	1501	GCNWSNYTNTYCAVGTNWSNAAAYGAYTAYWSNTTYATHMGNTAYTAYACNMGNCNATH	1560
Qy	521	TyrGlnPheGlnPheGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis	540
Db	1561	TAYCARTTYCARTTYCARGARCCNTNTTYGARGCNCNAAATYAAAYGWSNYSNCAAY	1620
Qy	541	LysCysAspIleSerAsnSerThrGluAlaGlyGlnLysLeuPheAsnMetLeuArgLeu	560
Db	1621	AARTCYGAYATHWSNAAYSNACNARGCNGCNCARAAAYTYTNAARATGYTNWSNYTN	1680
Qy	561	GlyLysSerGluProTrpThrLeuAlaLeuGluAsnValValGlyAlaLysAsnMetAsn	580
Db	1681	GGNAAYSNGARCCNTGGACNAAARGCNTNGARAAYGTNGTNGGNCNMGNAAYATGGAY	1740
Qy	581	ValArgProLeuLeuAsnTyrPheGluProLeuPheThrTrpLeuLysAspGlnAsnLys	600
Db	1741	GTNAAARCCNTNYTNAAYTAYTTYCARGCNTNTTYGYTGYTGGYTNAAARGCARAAAYMG	1800
Qy	601	AsnSerPheValGlyTrpSerThrAspTrpSerProTyrAlaAspGlnSerIleLysVal	620
Db	1801	AAYSNTTYTGTTGGTGGAAACNARGTGGWSNCCNTAYCGNGAYCARWSNATHAARGTN	1860
Qy	621	ArgIleSerLeuLysSerAlaLeuGlyAspLysAlaTyrGluTrpAsnAspAsnGluMet	640
Db	1861	MGNATHWSYTNAAARWSNCCNYTNGNCCNAAAYGCNTAYGARTGGACNAAAYAGARATG	1920
Qy	641	TyrLeuPheArgSerSerValAlaTyrAlaMetArgGlnTyrPheLeuLysValLysAsn	660
Db	1921	TTYTYTNTTYMGNWSNWSNTGCCNTAYGCNATGMCNAAARTAYTYWSNATHATHAARAY	1980
Qy	661	GlnMetIleLeuPheGlyGluGluAspValArgValAlaAsnLeuLysProArgIleSer	680
Db	1981	CARACNTGCCNTTYTNGARGARGAYGTNMGNGTWSNGAYTYTNAARCCNMGNTNWSN	2040
Qy	681	PheAsnPhePheValThrAlaProLysAsnValSerAspIleIleProArgThrGluVal	700
Db	2041	TTYTAYTTYTYTGNAACNWSNCCNAAAYGTNWSNGAYTGNATHCCNMGNWSNGARGTN	2100
Qy	701	GluLysAlaIleArgMetSerArgIleAsnAspAlaPheArgLeuAsnAspAsn	720
Db	2101	GARGAYCGNATHMGNATGWSNMGNGNMGNATHAAYGAYGTNTTYGGNYTNAAYGAYRAY	2160
Qy	721	SerLeuGluPheLeuGlyIleGlnProThrLeuGlyProProAsnGlnProValSer	740
Db	2161	WSNYTNGARTTYTNGGNATHCAYCCNACNYTNGARCCNCCNTAYCARCCNCCNGTNACN	2220
Qy	741	IleTrpLeuIleValPheGlyValIleValMetGlyValIleValValGlyValIleLeu	760

DB	2221	ATHGGYTNATHATHTTTCGGTNGTGNATGCGNTNGTNGTNGTNGGNATHATHATHYT	2280
QY	761	IlePheThrGlyIleArgAspArgLysLysAsnLysAlaArgSerGlyGluAsnPro	780
DB	2281	ATHGTNACGGGNATHAARGCGNMGNAARAARAARAAYGARACNAARMGNGARGAAYCCN	2340
QY	781	TyrAlaSerIleAspIleSerLysGlyGluAsnAsnProGlyPheGlnAsnThrAspAsp	800
DB	2341	TAYGAYWSNATGGATATGCGNARGNGARGWSNAAAYCGCGGNTTTCARAAAYWSNGAYGAY	2400
QY	801	ValGlnThrSerPhe	805
DB	2401	GCNCARACNWSNTTY	2415
RESULT 14			
ID	AAQ10328	standard; DNA; 2477 BP.	
AC	AAQ10328;		
DT	10-APR-1991	(first entry)	
DE	Encodes human testicular angiotensin conversion enzyme.		
KW	human testicular angiotensin conversion enzyme; tACE;		
KW	male sterility; ss.		
OS	Homo sapiens.		
FH	Key	Location/Qualifiers	
FT	CDS	29..2227	
FT		/*tag= a	
FT		/product= human tACE	
FT	Peptide	29..91	
FT		/*tag= b	
FT		/label= signal peptide	
XX	WO9100354-A.		
PN	10-JAN-1991.		
PD			
XX	05-JUL-1990;	90WO-FR00513.	
XX	05-JUL-1989;	89FR-0009062.	
XX	{INRM }	INST NAT SANTE RECH.	
PA	Soubrier F, Alhenc-Gelas F, Hubert C, Corvol P;		
XX			
XX	WPI: 1991-036748/05.		
DR	P-PSDB; AARI0426.		
XX			
PT	Nucleic acid - encoding human testicular angiotensin conversion		
PT	enzyme, used e.g. for in vitro detection of enzyme in organism		
XX			
PS	Claim 1; Fig 1: 48pp; French.		
XX			
CC	A bank of human testicular cDNA in Lambda gtl1 was screened with a		
CC	probe containing the final 3248 nucleotides of endothelial ACE. The		
CC	complete sequence of tACE was reconstructed from 4 separate clones.		
CC	It encodes a 711 amino acid mature protein and a 21 residue signal		
CC	peptide. The 228-2224 sequence is identical to the 1944-3940 region		
CC	of endothelial ACE. The isolated nucleic acid sequence is inserted		
CC	into a plasmid for expression of polypeptides. The invention also		
CC	covers parts of the sequence comprising all or part of the 29-229		
CC	sequence, any sequence differing from tACE only by silent		
CC	substitutions and nucleic acids which hybridise to tACE.		
XX			
SQ	Sequence 2477 BP; 536 A; 811 C; 695 G; 435 T; 0 other;		

Alignment Scores:

Pred. No.: 6.37e-121 Length: 2477

Score: 1344.00 Matches: 259  
Percent Similarity: 60.97% Conservative: 119  
Best Local Similarity: 41.77% Mismatches: 204  
Query Match: 31.32% Indels: 38  
DB: 12 Gaps: 10

US-09-635-501-2 (1-805) x AAQ10328 (1-2477)

QY 15 ThrAlaAlaGlnSer-----ThrIleGluGluGlnAlaLysThrPheLeuAsp 30  
DB 209 ACATCGCCAGAGCCCAAACTGGTACTGATGAGGTGGAGCCAGCAAGTTTGTGGAG 268  
QY 31 LysPheAsnHisGluAlaGluAspPheTyrglnSerSerLeuAlaSerTrpAsnTyr 50  
DB 269 GAATATGACCGGACATCCAGGTGGTGGAAAGAGATGTCGCGAGGCCAACCTGGAATAC 328  
QY 51 AsnThrAsnIleThrGluGlu-----AsnValGlnAsnMet 62  
DB 329 AACACCAACATCACCAGAGACCAGCAAGATTCTGTCGACAGAACATGCAAAATAGCC 388  
QY 63 AsnAsnAlaGlyAspLysTrpSerAlaPheLeuLysGluGlnSerThrLeuAlaGlnMet 82  
DB 389 AACCAACACC-----CTGAAGTACGGCACCCAGCCAGGAG 424  
QY 83 TyrProLeuGlnGluIleGlnAsnLeuThrValLysLeuGlnLeuGlnAlaLeuGlnGln 102  
DB 425 TTTGATGTGAACCAAGTTGCAGAACCACTATCAAGCGGATCATAAAGAGGTTTCAGGAC 484  
QY 103 AsnGlySerSerValLeuSerGluAspLysSerLysArgLeuAsnThrIleLeuAsnThr 122  
DB 485 CTGAAGCGGCGGCTCCCTGCCAGGAGCTGGAGGAGTACAAAGATCCTGTGGAT 544  
QY 123 MetSerThrIleTyrSerThrGlyLysValCysAsnProAspAsnProGlnGluCysLeu 142  
DB 545 ATGGAACCACTACAGCGTGGCCACTGTGTGCCACCGCAATGGC-----AGCTGCGCTG 598  
QY 143 LeuLeuGluProGlyLeuAsnGluIleMetAlaAsnSerLeuAspTyrAsnGluArgLeu 162  
DB 599 CAGCTCGAGCCAGATCTGACCAATGTGATGGCCACATCCCGGAAATATGAAGACCTGTTA 658  
QY 163 TrpAlaTrpGluSerTrpArgSerGluValGlyLysGlnLeuArgProLeuTyrGluGlu 182  
DB 659 TGGCGATGGGAGGGCTGGGAGACAGAGGGGGAGAGCCATCTCCAGTTTATCCCGAAA 718  
QY 183 TyrValValLeuLysAsnGluMetAlaArgAlaAsnHisTyrGluAspTyrGlyAspTyr 202  
DB 719 TACGTGGAACTCATCAACAGCGCTCCCGCTCAATGGCTATGATGATGAGGGGACTCG 778  
QY 203 TrpArgGlyAspTyrGluValAsnGlyValAspGlyTyrAspTyrSerArgGlyGlnLeu 222  
DB 779 TGGAGGTCTATGTACGAGACACCATCCCTGGAG----- 811  
QY 223 IleGluAspValGluHisThrPheGluGluIleLysProLeuTyrGluHisLeuHisAla 242  
DB 812 ---CAAGACCTGGAGGGCTCTTCCAGGAGCTTCCAGGAGCTTACCTCAACCTGCATGCC 868  
QY 243 TyrValArgAlaLysLeuMetAsnAlaTyr---ProSerTyrIleSerProIleGlyCys 261  
DB 869 TACGTGGCGCGGCGCTGACCGCTACTACGGGGCCGACGACATCAACCTGGAGGGGCC 928  
QY 262 LeuProAlaHisLeuLeuGlyAspMetTrpGlyArgPheTrpThrAsnLeuTyrSerLeu 281  
DB 929 ATTCTGTCTACCTGCTGGGAAACATGTGGCGCAGACCTGCTGCTCAACATCTATGACTTG 988  
QY 282 ThrValProPheGlyGlnLysProAsnIleAspValThrAspAlaMetValAspGlnAla 301  
DB 989 GTGGTGCCTTCCCTTACGCGCCCTCGATGGACACCAAGAGGCTATGCTAAAGCAGGGC 1048  
QY 302 TrpAspAlaGlnArgIlePheLysGluAlaGluLysPheValSerValGlyLeuPro 321  
DB 1049 TGGACGCCAGGAGGATGTTTAAGGAGGCTGATGTTCTTACCTCCCTGGGCTGCTG 1108  
QY 322 AsnMetThrGlnGlyPheTrpGluAsnSerMetLeuThrAspProGlyAsnValGlnLys 341

DB 1109 CCCGTGCTCCTGAGTTCTTGGAAACAAGTCGATGCTGGAGAAGCAACCCGAGCGGGGAG 1168  
QY 342 AlaValCysHisProThrAlaTrpAspLeuGlyLysGly---AspPheArgIleLeuMet 360  
DB 1169 GTGGTCTCCACCGCTCGGCTGGGACTTCTCAACGGCAAGGACTTCCGGATCAACGAG 1228  
QY 361 CysThrLysValThrMetAspPheLeuThrAlaHisHisGluMetGlyHisIleGln 380  
DB 1229 TGCACCAACCGTGAACCTGGAGGACTGGTGGTGGCCACCACCAAAATGGGCCACATCCAG 1288  
QY 381 TyrAspMetAlaTyrAlaAlaGlnProPheLeuLeuArgAsnGlyAlaAsnGluGlyPhe 400  
DB 1289 TATTTCATGTCAGTACAAAGACTTACCTGTGGCTTGGAGGAGGTGCCAACCCCGGCTTC 1348  
QY 401 HisGluAlaValGlyLysLeuMetSerLeuSerAlaAlaThrProLysHisLeuLysSer 420  
DB 1349 CATGAGGCCATTTGGGAGCTGTAGCCCTCTAGTGTCTAGCCCAAGCAGCTGCACAGT 1408  
QY 421 IleGlyLeuLeuSerProAspPheGlnGluAspAsnGluThrGluIleAsnPheLeuLeu 440  
DB 1409 CTCACCTGCTGAGCAGTGGGTGGCAGGAC---GAGCATGACATCACTTCTGATG 1465  
QY 441 LysGlnAlaLeuThrIleValGlyThrLeuProPheThrTyrMetLeuGluLysTrpArg 460  
DB 1466 AGGTGGCCCTTGACAAGATCGCCTTTATCCCTTACGTACCTCGTGCATCAGTGGGC 1525  
QY 461 TrpMetValPheLysGlyGluIleProLysAspGlnTrpMetLysLysTrpTrpGluMet 480  
DB 1526 TGGAGGGTATTTGATGGAGCATCACCAGGAGAACTATAACAGGAGTGGTGGAGCCCTC 1585  
QY 481 LysArgGluIleValGlyValGluProValProHisAspGluThrTyrCysAspPro 500  
DB 1586 AGGCTGAAGTACAGGGCTCTGCCCCCAAGTGGCCAGGACTCAAGGTGACTTTGACCCA 1645  
QY 501 AlaSerLeuPheHisValSerAsnAspTyrSerPheIleArgTyrTyrThrArgThrLeu 520  
DB 1646 GGGGCCAAGTTCACATCTCTTACGCGTGGCTTACATCAGGTACTTTGTACGTTCAATC 1705  
QY 521 TyrGlnPheGlnPheGlnGluAlaLeuCysGlnAlaAlaLysHisGluGlyProLeuHis 540  
DB 1706 ATCCAGTTCAGTTCACAGGAGGACTGTGCCAGGAGCTGGCCACACGCGCCCTGAC 1765  
QY 541 LysCysAspIleSerAsnSerThrGluAlaGlyGlnLysLeuPheAsnMetLeuArgLeu 560  
DB 1766 AAGTGTGACATCTACCAAGTCCAAAGGCGGGGAGCCCTGGCCACCCCGCATGAAGCTG 1825  
QY 561 GlyLysSerGluProTrpThrLeuAlaLeuGluAsnValValGlyAlaLysAsnMetAsn 580  
DB 1826 GGCTTACAGTGGCGGCGGAGCCATGACGATGATCAGGGCCAGCCCAACATGAGC 1885  
QY 581 ValArgProLeuLeuAsnTyrPheGluProLeuPheThrTrpLeuLysAspGlnAsnLys 600  
DB 1886 GCCTCGGCCATGTGACTACTTCAAGCGCTGTGTGGACTGGCTCCGACGAGAGACGAG 1945  
QY 601 -----AsnSerPheValGlyTrp---SerThrAspTrpSerProTyrAlaAspGlnSer 617  
DB 1946 CTGCATGGGAGAGAGCTGGGCTGGCCGAGTACACTGACGCGCCGAACCTCCGCTCGCTCA 2005

RESULT 15  
AAA38330  
ID AAA38330 standard; DNA; 4020 BP.  
XX  
AC AAA38330;  
XX  
DT 21-AUG-2000 (first entry)  
XX  
DE Human angiotensin-converting enzyme (ACE) coding region.  
XX  
KW Angiotensin-converting enzyme gene; ACE; coding region; polymorphism;  
KW polymorphic marker; cardiovascular disease; myocardial infarction;  
KW unstable angina; hypertension; atherosclerosis; stroke; prognosis;  
KW drug screening; treatment outcome; human; ds.

XX OS Homo sapiens.  
XX PN WO200022166-A2.  
XX PD 20-APR-2000.  
XX PF 13-OCT-1999; 99WO-1801678.  
XX PR 14-OCT-1998; 98US-0104286.  
XX PR 14-OCT-1998; 98US-0104302.  
XX PA (EURO-) EURONA MEDICAL AB.  
XX PI Norberg LT, Andersson MK, Lindstrom PHR, Jonsson L;  
XX WPI; 2000-318010/27.  
XX  
PT Assessing cardiovascular status in humans involves comparing test  
PT polymorphic pattern comprising polymorphic positions within genes  
PT encoding specific proteins, with reference polymorphic pattern -  
XX  
PS Disclosure: Page 114-115; 126pp; English.  
XX  
CC The invention relates to a novel method of assessing the cardiovascular  
CC status in an individual and to newly identified polymorphisms in the  
CC genes encoding angiotensin-converting enzyme (ACE), angiotensin II  
CC receptor type 1 (AT1) and type 2 (AT2), angiotensinogen (AGT), renin,  
CC aldosterone synthase, endothelin receptor type A and beta-adrenergic  
CC receptors 1 and 2. The method comprises determining the sequence at one  
CC or more polymorphic positions within these genes, and comparing the  
CC pattern of polymorphisms from the individual with a reference polymorphic  
CC pattern obtained from a population of individuals exhibiting a  
CC predetermined cardiovascular disease status. The polymorphic markers are  
CC useful for determining the predisposition of an individual to  
CC cardiovascular disorders such as myocardial infarction, unstable angina,  
CC hypertension, atherosclerosis and stroke. They are also useful for  
CC predicting the likely cardiovascular status of a patient given a  
CC treatment regimen comprising administration of cardiovascular drugs  
CC (e.g., ACE inhibitors, beta-adrenergic receptor antagonists (beta-  
CC blockers) or calcium channel blockers). One or more polymorphic markers  
CC provides a basis for predicting the outcome of a treatment regimen.  
CC Fragments of the genes comprising a polymorphic site may be used as  
CC primers and probes for detecting genetic polymorphisms or in molecular  
CC library arrays for high throughput screening. The genes, and the proteins  
CC they encode are useful in the screening of potential cardiovascular  
CC drugs. Determination of an individual's polymorphic pattern reduces or  
CC eliminates trial and error in selecting a treatment for a particular  
CC individual cardiovascular patient. It also provides the ability to  
CC eliminate patients from clinical trials who are predicted to be  
CC non-responsive, or at a risk for an adverse response, to a particular  
CC treatment regimen. Adverse results in an early trial can be evaluated to  
CC identify polymorphic patterns so that the adverse results can be  
CC correlated with a sub-population of the test population, permitting  
CC exclusion of such sub-populations from the treatment group. Beneficial  
CC drugs can be approved for use in the appropriate population, thereby  
CC decreasing the number of patients required for a clinical trial, which in  
CC turn decreases the duration and cost of such trials. Sequences A3828 and  
CC A38330 represent, respectively, intron 16 and the coding region of  
CC the human ACE gene (Genbank X62855, J04144). The polymorphic sites  
CC identified are 375A/C, 582C/T, 731A/G, 1060G/A, 1215C/T, 2193G/A,  
CC 2328A/G, 2741G/T, 3132C/T, 3387T/C, 3503G/C, 3906G/A; and a deletion of  
CC nucleotides 1451-1783 in intron 16.  
XX  
SQ Sequence 4020 BP; 857 A; 1261 C; 1174 G; 728 T; 0 other;

Alignment Scores:  
Pred. No.: 6,14e-120 Length: 4020  
Score: 1337.00 Matches: 255  
Percent Similarity: 61.05% Conservative: 118  
Best Local Similarity: 41.73% Mismatches: 204  
Query Match: 31.16% Indels: 34  
DB: 21 Gaps: 9

US-09-635-501-2 (1-805) x AAA38330 (1-4020)  
QY 20 ThrIleGluGluGlnAlaLysThrPheLeuAspLysPheAsnHisGluAlaGluAspLeu 39  
DB 1952 ACTGATGAGGCTGAGGCCAGCAAGTTTGTGAGGAATATGACCGACATCCAGGTGGTG 2011  
QY 40 PheTyrGlnSerSerLeuAlaSerTrpAsnTyrAsnThrAsnIleThrGluGlu----- 57  
DB 2012 TGGAACGAGTATGCCGAGGCCAACTGGAACTACCAACACCAACATCACACAGAGACCAGC 2071  
QY 58 -----AsnValGlnAsnMetAsnAlaGlyAspLysTrpSerAla 71  
DB 2072 AAGATTCTGCTGCAGAGAACATGCAATAGCCAAACACACACC----- 2113  
QY 72 PheLeuLysGluGlnSerThrLeuAlaGlnMetTyrProLeuGlnGluIleGlnAsnLeu 91  
DB 2114 -----CTGAAGTACGGCACCAGCCAGGAGGTTTGTGATGTGAACCACTTGCAGACACACC 2167  
QY 92 ThrValLysLeuGlnLeuGlnAlaLeuGlnAlaAsnGlySerSerValLeuSerGluAsp 111  
DB 2168 ACTATCAAGCGGATCATAAAGAGGTTTCAGGACCTAGAACGGGCGGCTGCTGCCCCAG 2227  
QY 112 LysSerLysArgLeuAsnThrIleLeuAsnThrMetSerThrIleTyrSerThrGlyLys 131  
DB 2228 GAGCTGGAGGAGTACAAACAGATCTCTGTTGGATATGAAACCCACCTACAGGCTGCCACT 2287  
QY 132 ValCysAsnProAspAsnProGlnGluCysLeuLeuLeuGluProGlyLeuAsnGluIle 151  
DB 2288 GTGCGCCACCCGAATGGC-----ACCTGCTGCAGCTCGAGCCAGATCTCAGCAATGTG 2341  
QY 152 MetAlaAsnSerLeuAspTyrAsnGluArgLeuTrpAlaTrpGluSerTrpArgSerGlu 171  
DB 2342 ATGGCCACATCCCGGAAATATGAAGACCTTATGGCATGGGAGGCTGGCGAGACACAG 2401  
QY 172 ValGlyLysGlnLeuArgProLeuTyrGluGluTyrValValLeuLysAsnGluMetAla 191  
DB 2402 GCGGGAGAGGCATCTCCAGTTTACCCGAAATACGTGGAACCTCATCAACGAGCTGCC 2461  
QY 192 ArgAlaAsnHisTyrGluAspTyrGlyAspTyrTrpArgGlyAspTyrGluValAsnGly 211  
DB 2462 CGGCTCAATGGCTATGTAGATCGAGGGGACTCGTGGAGGTCTATGTACGAGACACCATCC 2521  
QY 212 ValAspGlyTyrAspTyrSerArgGlyGlnLeuIleGluAspValGluHisThrPheGlu 231  
DB 2522 CTGGAG-----CAAGACCTGGAGCGGCTCTTCCAG 2551  
QY 232 GluLeLysProLeuTyrGluHisLeuHisAlaTyrValArgAlaLysLeuMetAsnAla 251  
DB 2552 GAGCTGCAGCCACTCTACCTCAACCTGCATGCCTACGTGCGCGCGGCGCTGCACCGTCAC 2611  
QY 252 Tyr---ProSerTyrIleSerProIleGlyCysLeuProAlaHisLeuLeuGlyAspMet 270  
DB 2612 TACGGGGCCAGCACATCAACCTGGAGGGGCCCATCTCTCTCCTCACCTGCTGGGGAACATG 2671  
QY 271 TrpGlyArgPheTrpThrAsnLeuTyrSerLeuThrValProPheGlyGlnLysProAsn 290  
DB 2672 TGGCGCGAGACCTGGTCCCAACATCTATGACTTGGTGGTGGCTTCCCTTTCAGCGCCCTCG 2731  
QY 291 IleAspValThrAspAlaMetValAspGlnAlaTrpAspAlaGlnArgIlePheLysGlu 310  
DB 2732 ATGGACACCACAGAGGCTATGCTAAAGCAGGCGCTGGAGGCCAGGAGGATGTTTAAAGGAG 2791  
QY 311 AlaGluLysPhePheValSerValGlyLeuProAsnMetThrGlnGlyTrpGluAsn 330  
DB 2792 GCTGATGATTTCTCACCTCCCTGGGGCTGCTGCCCGTGCCTCCTGAGTTCCTGGAAACAG 2851  
QY 331 SerMetLeuThrAspProGlyAsnValGlnLysAlaValCysHisProThrAlaTrpAsp 350  
DB 2852 TCGATGCTGGAGAGCCCAACCGACGGCGGAGGTTGTCTGCCACCGCTCGGCTGGGAC 2911  
QY 351 LeuGlyLysGly---AspPheArgIleLeuMetCysThrLysValThrMetAspAspPhe 369

Db 2912 TTCTACAAGCGCAAGGACTTCGGGATCAAGCACTGCACCACCGCTGAACCTTGGAGGACCTG 2971  
Qy 370 LeuThrAlaHisHisGluMetGlyHisIleGlnTyrAspMetAlaTyrAlaAlaGlnPro 389  
Db 2972 GTGGTGGGCCACCAAGGATGGCCACATCCAGTATTTTCAGTACAGTACAAAGACTTACCT 3031  
Qy 350 PheLeuLeuArgAsnGlyAlaAsnGluGlyPheHisGluAlaValGlyGluIleMetSer 409  
Db 3032 GTGGCCTTGAGGGAGGTGCCAACCCCGGCTTCATGAGGCCATTTGGGACGTGCTAGCC 3091  
Qy 410 LeuSerAlaAlaThrProLysHisLeuLysSerIleGlyLeuLeuSerProAspPheGln 429  
Db 3092 CTCTCAGTCTACGCCCAAGCACTGCACAGTCTCAACCTGCTGAGCAGTGGGGTGGC 3151  
Qy 430 GluAspAsnGluThrGluIleAsnPheLeuLeuLysGlnAlaLeuThrIleValGlyThr 449  
Db 3152 AGCGAC--GAGCATGACATCAACTTCTGATGAAGATGGCCCTTGACAGATCGCCTTT 3208  
Qy 450 LeuProPheThrTyrMetLeuGluLysTyrArgTyrMetValPheLysGlyGluIlePro 469  
Db 3209 ATCCCTTCAGTACCTCGTCGATCATGCTGGCTGGAGGGTATTTGATGGAAGCATCAC 3268  
Qy 470 LysAspGluTrpMetLysLysTyrTrpGluMetLysArgGluIleValGlyValValGlu 489  
Db 3269 AAGGAGAACTATTAACAGGAGTGTGGAGCTCAGGCTCAGGCTGAAGTACCAGGCGCTCTGCCCC 3328  
Qy 490 ProValProHisAspGluThrTyrCysAspProAlaSerLeuPheHisValSerAsnAsp 509  
Db 3329 CCAGTCCCGCAGGACTCAAGTGACTTTGACCCAGGGGCCAAAGTTCACATTCCTTCTAGC 3388  
Qy 510 TyrSerPheIleArgTyrTyrThrArgThrLeuTyrGlnPheGlnPheGlnGluAlaLeu 529  
Db 3389 GTGCCCTTACATCAGGTACTTTGTGACGCTTCATCATCCAGTTCCAGTTCCACGAGGCAC 3448  
Qy 530 CysGlnAlaAlaLysHisGluGlyProLeuHisLysCysAspIleSerAsnSerThrGlu 549  
Db 3449 TGCCAGGACGTGGCCACAGCGGGCCCCCTGCACAGTGTGACATCTACCAGTCCAAAGGAG 3508  
Qy 550 AlaGlyGlnLysLeuPheAsnMetLeuArgLeuGlyLysSerGluProTrpThrLeuAla 569  
Db 3509 GCCGGCAGCGCCTGGCGACCGCCATGAAGCTGGGCTTCAGTAGGCCGTGGCGGGAAGCC 3568  
Qy 570 LeuGluAsnValValGlyAlaLysAsnMetAsnValArgProLeuLeuAsnTyrPheGlu 589  
Db 3569 ATGCAGCTGATCAGGGGCCAGCCCAACATGAGCGCTCGGCCCATGTTGAGCTACTTCAAG 3628  
Qy 590 ProLeuPheThrTrpLeuLysAspGlnAsnLys-----AsnSerPheValGlyTrp--- 606  
Db 3629 CCGCTGCTGACTGGCTCCGACCGGAGAACGAGTGCATGGGAGAGCTGGGCTGGCCG 3688  
Qy 607 SerThrAspTrpSerProTyrAlaAspGlnSer 617  
Db 3689 CAGTACAACGTGGAGCGCGCACTCCGCTCGCTCA 3721

Search completed: October 9, 2002, 18:07:43  
Job time : 295 secs